

Result No.	Score	Query		Length	DB	ID	Description
		Match					
1	943	99.0	3011	1	POLG_HCV1		P26664 h genome po
2	927	97.3	3011	1	POLG_HCV1		P27958 h genome po
3	852	93.6	3010	1	POLG_HCVTW		P29846 h genome po
4	891	93.5	3010	1	POLG_HCVBK		P26663 h genome po
5	887	93.1	3010	1	POLG_HCV1A		P26662 h genome po
6	884	92.8	3010	1	POLG_HCVJT		Q00269 h genome po
7	714	74.9	3033	1	POLG_HCVJ8		P26661 h genome po
8	712	74.7	3033	1	POLG_HCVJ6		P26660 h genome po
9	87	9.1	209	1	PAAD_PSEAE		Q9h008 pseudomonas
10	84	8.8	321	1	HQOA_ARATH		Q9h067 arabidopsis
11	82	8.6	452	1	ANMP_HUMAN		Q9sel7 arabidopsis
12	82	8.6	485	1	Y136_TREPA		Q93683 homo sapien
13	80.5	8.4	437	1	DEG1_ARATH		O83172 treponema p
14	75	7.9	253	1	CAG3_BOVIN		O29609 arabidopsis
15	74.5	7.8	415	1	ZP3_RABIT		P05805 bos taurus
16	74.5	7.8	776	1	HYPE_AZOVI		P48833 oryctolagus
17	74.5	7.8	911	1	TB11_NEJMB		P40596 azotobacter
18	74	7.8	326	1	PANE_RH10O		Q09056 neisseria m
19	73.5	7.7	263	1	GRAX_MOUSE		Q96785 rhizobium l
20	73	7.7	730	1	HELS_METNA		C35295 mus musculu
21	72.5	7.6	257	1	GRAM_HUMAN		Q8p2r7 methanosarc
22	72	7.6	627	1	SAD1_MOUSE		P51124 homo sapien
23	72	7.6	1527	1	CATH_MOUSE		Q60710 mus musculu
24	72	7.6	2663	1	CENE_HUMAN		P38061 mus musculu
25	72	7.6	3491	1	ERV1_SACER		Q02224 homo sapien
26	71.5	7.5	248	1	GRAD_MOUSE		Q01131 saccharopol
27	71.5	7.5	323	1	VPRT_SMRW		P11033 mus musculu
28	71	7.5	219	1	SPR1_IPOBA		P21407 squirrel sc
29	71	7.5	336	1	UL16_EBV		P14715 ibomoea bat
30	71	7.5	328	1	PG12_RALSO		P03221 epstein-bar
31	71	7.5	1280	1	ITR1_RAT		P20041 raistonia s
32	70.5	7.4	264	1	CTRL_HUMAN		P18614 rattus norv
33	70.5	7.4	659	1	V572_HEVME		P40313 homo sapien
34	70.5	7.4	659	1	V572_HEVME		Q03500 hepacillus g

InterPro: IPR002521; HCV_core.
 DR InterPro: IPR002519; HCV_env.
 DR InterPro: IPR002531; HCV_NS1.
 DR InterPro: IPR002518; HCV_NS2.
 DR InterPro: IPR004109; HCV_NS3.
 DR InterPro: IPR006745; HCV_NS4a.
 DR InterPro: IPR001490; HCV_NS4b.
 DR InterPro: IPR002868; HCV_NS5a.
 DR InterPro: IPR002166; HCV_RdRp.
 DR InterPro: IPR001650; Helicase_C.
 DR InterPro: IPR007095; RNA_pol_DS_PS.
 DR InterPro: IPR007094; RNA_pol_PSVir.
 DR Pfam: PF01543; HCV_capsid; 1.
 DR Pfam: PF01542; HCV_core; 1.
 DR Pfam: PF01539; HCV_env; 1.
 DR Pfam: PF01560; HCV_NS1; 1.
 DR Pfam: PF01538; HCV_NS2; 1.
 DR Pfam: PF02907; HCV_NS3; 1.
 DR Pfam: PF01006; HCV_NS4a; 1.
 DR Pfam: PF01001; HCV_NS4b; 1.
 DR Pfam: PF01506; HCV_NS5a; 1.
 DR Pfam: PF00271; helicase_C; 1.
 DR Pfam: PF00398; Viral_RdRp; 1.
 DR ProDom: PD086062; HCV_NS1; 1.
 DR SMART: SM00487; DEXDC; 1.
 DR PolyProtein: Glycoprotein; Transferase; RNA-directed RNA polymerase;
 KW Core protein; Coat protein; Envelope protein; Helicase; ATP-binding;
 KW Transmembrane; Nonstructural protein; Hydrolase; Serine protease;
 KW 3D-structure.
 FT INIT_MET 1 1 REMOVED FROM CAPSID PROTEIN C BY THE
 CELLULAR AMINOPEPTIDASE.
 FT CHAIN 1 115
 FT CHAIN 116 191
 FT CHAIN 192 383
 FT CHAIN 384 729
 FT CHAIN 730 1006
 FT CHAIN 1007 1515
 FT CHAIN 1616 1862
 FT CHAIN 1863 2013
 FT CHAIN 2014 3011
 FT CHAIN 3011 359
 FT TRANSMEM 347 359
 FT ACT_SITE 1083 1083
 FT ACT_SITE 1137 1137
 FT ACT_SITE 1165 1165
 FT NP_BIND 1230 1237
 FT SITE 1316 1319
 FT CARBOHYD 196 196
 FT CARBOHYD 209 209
 FT CARBOHYD 234 234
 FT CARBOHYD 305 305
 FT CARBOHYD 417 417
 FT CARBOHYD 423 423
 FT CARBOHYD 430 430
 FT CARBOHYD 448 448
 FT CARBOHYD 476 476
 FT CARBOHYD 532 532
 FT CARBOHYD 540 540
 FT CARBOHYD 556 556
 FT CARBOHYD 576 576
 FT CARBOHYD 623 623
 FT CARBOHYD 645 645
 FT CARBOHYD 2041 2041
 FT CARBOHYD 2077 2077
 FT CARBOHYD 2240 2240
 FT CARBOHYD 2364 2364
 FT CARBOHYD 2789 2789
 SQ SEQUENCE 3011 AA; 327197 MW; 65F8C9447FCE5AF9 CRC64;
 Query Match 99.0%; Score 943; Db 1; Length 3011;
 Best Local Similarity 98.4%; Pred. No. 3.4e-82;
 Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

Db 1026 LAPITAYAAQTGRLGCIITSLTGRKNOVEQVIVSTAAQTFLATCINGVCWTVYHGA 1085
 QY 61 GTRTIASPKGPVIQMYTIVDKOLVGWPAPOGSKSLPTCGSSDIYIVTRHADVIPVRRR 120
 Db 1086 GTRTIASPKGPVIQMYTIVDKOLVGWPAPOGSKSLPTCGSSDIYIVTRHADVIPVRRR 1145
 QY 121 GDSRGLSLSPRISYLVKSGSGGGLLCPAGHAGIFRAAVCTRGVAKAVDFIPVESLETTM 180
 Db 1146 GDSRGLSLSPRISYLVKSGSGGGLLCPAGHAGIFRAAVCTRGVAKAVDFIPVENLETTM 1205
 QY 181 RS 182
 Db 1206 RS 1207
 RESULT 2
 POLG_HCVH STANDARD; PRT; 3011 AA.
 AC P27958;
 DT 01-AUG-1992 (Rel. 23, Created)
 DT 01-AUG-1992 (Rel. 23, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Genome polyprotein [Contains: Capsid protein C (Core protein) (P22);
 DE Envelope glycoprotein E1 (GP32) (GP35); Envelope glycoprotein E2
 DE (GP68) (GP70) (NS1); protein P7; Nonstructural protein NS2 (P21)
 DE (EC 3.4.99.-); Protease/helicase NS3 (P70) (Hepacivirin)
 DE (EC 3.4.21.98); Nonstructural protein NS4A (P4); Nonstructural protein
 DE NS4B (P27); Nonstructural protein NS5A (P56); Nonstructural protein
 DE NS5B (P66) (P70) (RNA-directed RNA polymerase) (EC 2.7.7.48)].
 OS Hepatitis C virus (isolate H) (HCV).
 OC Viruses; ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
 OC Hepacivirus.
 OX NCB:TaxID=11106;
 RN [1]-TaxID=11106;
 SEQUENCE FROM N.A.
 RP MEDLINE=92052256; PubMed=1658800;
 RA Inchauspe G., Zebedee S., Lee D.H., Sugitani M., Nasoff M.,
 RA Pirace A.M.;
 RI "Genomic structure of the human prototype strain H of hepatitis C
 RI virus: comparison with American and Japanese isolates.";
 RL Proc. Natl. Acad. Sci. U.S.A. 88:10292-10296(1991).
 RN [2];
 RP X-RAY CRYSTALLOGRAPHY (2.1 ANGSTROMS) OF 1207-1657.
 RX MEDLINE=97331322; PubMed=9187654;
 RA Yao N., Hesson T., Cable M., Hong Z., Kwong A.D., Le H.V., Weber P.C.;
 RL "Structure of the hepatitis C virus RNA helicase domain.";
 RN Nat. Struct. Biol. 4:463-467(1997).
 RN [3];
 RP X-RAY CRYSTALLOGRAPHY (2.2 ANGSTROMS) OF 1192-1657.
 RX MEDLINE=9815432; PubMed=9493270;
 RA Kim J.L., Morgenstern K.A., Griffith J.P., Dwyer M.D., Thomson J.A.,
 RA Murcko M.A., Lin C., Caron P.R.;
 RI "Hepatitis C virus NS3 RNA helicase domain with a bound
 RI oligonucleotide: the crystal structure provides insights into the mode
 RI of unwinding.";
 RL Structure 6:89-100(1998).
 CC -!- FUNCTION: PROTEASE NS2 IS RESPONSIBLE FOR THE CLEAVAGE OF NS2-NS3.
 CC -!- FUNCTION: PROTEASE NS3 IS RESPONSIBLE FOR THE CLEAVAGE OF
 CC NS3-NS4A, NS4A-NS4B, NS4B-NS5A AND NS5A-NS5B.
 CC -!- FUNCTION: NS4A FORMS A COMPLEX WITH NS3 AND IS ESSENTIAL FOR THE
 CC ACTIVATION OF NS3.
 CC -!- FUNCTION: NS5A SEEMS TO HAVE A TRANSCRIPTIONAL ACTIVATORY ROLE.
 CC -!- FUNCTION: NS5B IS A RNA-DEPENDENT RNA POLYMERASE THAT PLAYS AN
 CC ESSENTIAL ROLE IN THE VIRUS REPLICATION.
 CC -!- CATALYTIC ACTIVITY: Hydrolysis of four peptide bonds in the viral
 CC precursor polyprotein, commonly with Asp or Glu in the p6
 CC position, Cys or Thr in P1 and Ser or Ala in P1'.
 CC -!- CATALYTIC ACTIVITY: N nucleoside triphosphate = N diphosphate +
 CC [RNA](N).
 CC -!- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
 CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS: E1
 CC AND E2. THE NUCLEOCAPSID IS A COMPLEX OF PROTEIN C AND MRNA.

CC -!- PTM: THE STRUCTURAL PROTEINS C, E1 AND E2 ARE PRODUCED BY
 CC PROTEOLYTIC PROCESSING BY THE HOST SIGNAL PEPTIDASES.
 CC -!- SIMILARITY: THE NS2 PROTEASE BELONGS TO PEPTIDASE FAMILY U39.
 CC -!- SIMILARITY: THE NS3 PROTEASE BELONGS TO PEPTIDASE FAMILY S29.
 CC -----
 CC THIS SWISS-PROT entry is copyright. It is produced through a co-laboration
 CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See [http://www.isb-sib.ch/](http://www.isb-sib.ch/announce/)
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL: M67463; AAA45534.1; ..
 DR PIR: A36814; GNMVCH
 DR PDB: 1HEI; 25-NOV-98.
 DR PDB: 1AIV; 16-FEB-99.
 DR PDB: 1AIR; 17-JUN-98.
 DR MEROPS: S29.001; ..
 DR MEROPS: U39.001; ..
 DR TRANSFAC: T04155; ..
 DR InterPro: IPR00141C; DEAD.
 DR InterPro: IPR002522; HCV_capsid.
 DR InterPro: IPR002521; HCV_core.
 DR InterPro: IPR002519; HCV_env.
 DR InterPro: IPR002531; HCV_NS1.
 DR InterPro: IPR002518; HCV_NS2.
 DR InterPro: IPR004109; HCV_NS3.
 DR InterPro: IPR000745; HCV_NS4a.
 DR InterPro: IPR001490; HCV_NS4b.
 DR InterPro: IPR002868; HCV_NS5a.
 DR InterPro: IPR002166; HCV_RGRP.
 DR InterPro: IPR001650; Helicase_C.
 DR InterPro: IPR007095; RNA_pol_DS_PS.
 DR InterPro: IPR007094; RNA_pol_PS_vir.
 DR Pfam: PF01543; HCV_capsid; 1.
 DR Pfam: PF01542; HCV_core; 1.
 DR Pfam: PF01539; HCV_env; 1.
 DR Pfam: PF01560; HCV_NS1; 1.
 DR Pfam: PF01538; HCV_NS2; 1.
 DR Pfam: PF02907; HCV_NS3; 1.
 DR Pfam: PF01066; HCV_NS4a; 1.
 DR Pfam: PF01061; HCV_NS4b; 1.
 DR Pfam: PF01506; HCV_NS5a; 1.
 DR Pfam: PF00271; helicase_C; 1.
 DR Pfam: PF00998; Viral_RdRP; 1.
 DR ProDom: PD186062; HCV_NS1; 1.
 DR SMART; SM00487; DEXDC; 1.
 KW Polyprotein; Glycoprotein; Transferase; RNA-directed RNA polymerase;
 KW Core protein; Coat protein; Helicase; ATP-binding;
 KW Transmembrane; Nonstructural protein; Hydrolase; Serine protease;
 KW 3D-structure.
 FT INIT_MET 1 1 REMOVED FROM CAPSID PROTEIN C BY THE
 FT CHAIN 1 1 CELLULAR ANTIGENPEPTIDASE;
 FT CHAIN 192 383 CAPSID PROTEIN C.
 FT CHAIN 384 746 ENVELOPE GLYCOPROTEIN E1.
 FT CHAIN 747 809 ENVELOPE GLYCOPROTEIN E2.
 FT CHAIN 810 1326 PROTEIN P7.
 FT CHAIN 1027 1657 NONSTRUCTURAL PROTEIN NS2.
 FT CHAIN 1658 1711 PROTEASE/HELICASE NS3.
 FT CHAIN 1712 1972 NONSTRUCTURAL PROTEIN NS4a.
 FT CHAIN 1973 2420 NONSTRUCTURAL PROTEIN NS4b.
 FT CHAIN 2421 3011 NONSTRUCTURAL PROTEIN NS5a.
 FT CHAIN 347 369 NONSTRUCTURAL PROTEIN NS5b.
 FT ACT_SITE 1083 1083 POTENTIAL.
 FT ACT_SITE 1107 1107 CHARGE RELAY SYSTEM (BY SIMILARITY).
 FT ACT_SITE 1165 1165 CHARGE RELAY SYSTEM (BY SIMILARITY).
 FT NP_BIND 1230 1237 ATP (POTENTIAL).
 FT SITE 1316 1319 DECH_BOX.
 FT CARBOHYD 196 196 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 209 209 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 234 234 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CHAIN 1 191
 FT CHAIN 192 383
 FT CHAIN 384 746
 FT CHAIN 747 809
 FT CHAIN 810 1326
 FT CHAIN 1027 1657
 FT CHAIN 1658 1711
 FT CHAIN 1712 1972
 FT CHAIN 1973 2420
 FT CHAIN 2421 3011
 FT TRANSMEM 347 369
 FT ACT_SITE 1083 1083
 FT ACT_SITE 1107 1107
 FT ACT_SITE 1165 1165
 FT NP_BIND 1230 1237
 FT SITE 1316 1319
 FT CARBOHYD 196 196
 FT CARBOHYD 209 209
 FT CARBOHYD 234 234

FT CARBOHYD 305 305 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 417 417 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 423 423 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 430 430 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 448 448 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 476 476 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 532 532 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 540 540 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 556 556 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 576 576 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 623 623 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 645 645 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT STRAND 1224 1226
 FT TURN 1232 1233
 FT TURN 1236 1238
 FT HELIX 1239 1246
 FT TURN 1247 1248
 FT STRAND 1251 1255
 FT HELIX 1258 1271
 FT TURN 1272 1272
 FT STRAND 1277 1280
 FT TURN 1281 1282
 FT STRAND 1283 1285
 FT STRAND 1291 1295
 FT HELIX 1296 1301
 FT TURN 1302 1303
 FT STRAND 1312 1316
 FT TURN 1317 1319
 FT HELIX 1323 1335
 FT TURN 1336 1340
 FT STRAND 1343 1347
 FT TURN 1352 1353
 FT TURN 1360 1361
 FT STRAND 1362 1366
 FT STRAND 1368 1368
 FT TURN 1373 1375
 FT TURN 1376 1377
 FT STRAND 1378 1380
 FT HELIX 1382 1385
 FT STRAND 1389 1393
 FT HELIX 1397 1409
 FT TURN 1410 1411
 FT STRAND 1414 1417
 FT TURN 1419 1420
 FT STRAND 1432 1436
 FT TURN 1438 1439
 FT STRAND 1450 1453
 FT STRAND 1456 1463
 FT STRAND 1471 1478
 FT STRAND 1480 1480
 FT HELIX 1481 1488
 FT TURN 1489 1490
 FT STRAND 1497 1501
 FT STRAND 1507 1507
 FT STRAND 1511 1511
 FT HELIX 1514 1527
 FT HELIX 1532 1544
 FT STRAND 1550 1550
 FT HELIX 1555 1564
 FT STRAND 1570 1578
 FT TURN 1579 1580
 FT HELIX 1584 1597
 FT TURN 1598 1598
 FT HELIX 1606 1611
 FT TURN 1614 1618
 FT STRAND 1622 1623
 FT STRAND 1627 1627
 FT STRAND 1635 1636
 FT HELIX 1640 1652
 SQ SEQUENCE 3011 AA: 327142 MW: 772CBB29CCD94753 CRC64;
 Query Match 97.3%; Score 927; DB 1; Length 3011;
 Best Local Similarity 96.2%; Pred. No. 1.2e-80;

```
Matches 175: Conservative 4: Mismatches 3: Indels 0: Gaps 0:
QY 1 MAPITAYAOOTRLLGCIITISITGRKNOVEEVOIVTSAAOTELATCINGCWIVYHCA 60
DB 1026 LAPITAYAOOTRLLGCIITISITGRKNOVEEVOIVTSATOTFLATCINGCWIVYHCA 1085
QY 61 GRTIASPKSPVOTYTNVDKIDLVGPAHOGSRSLTPCICGSSDLYLVTRHADVIPVR 120
DB 1086 GRTIASPKSPVOTYTNVDKIDLVGPAHOGSRSLTPCICGSSDLYLVTRHADVIPVR 145
QY 121 GDSRGSLLSPRISYLYKSGSGFLPCPAGHAGVCFRAAVCTRGVAKAVDFEIPVESLETIM 180
DB 1146 GDSRGSLLSPRISYLYKSGSGFLPCPAGHAGVCFRAAVCTRGVAKAVDFEIPVENLETIM 1205
QY 181 RS 182
DB 1206 RS 1207

RESULT 3
POLG_HCVTW STANDARD: PRT: 3010 AA.
AC P29846:
DT 01-APR-1993 (Rel. 25, Created)
DT 01-APR-1993 (Rel. 25, Last sequence update)
DE Genome polyprotein [Contains: Capsid protein C (Core protein) (P22);
DE Envelope glycoprotein E1 (GP32) (GP35); Envelope glycoprotein E2
DE (GP68) (GP70) (NS1); Protein P7; Nonstructural protein NS2 (P21)
DE (EC 3.4.22.-); Protease/helicase NS3 (P70) (Hepacivirin)
DE (EC 3.4.21.98); Nonstructural protein NS4A (P4); Nonstructural protein
DE NS4B (P27); Nonstructural protein NS5A (P56); Nonstructural protein
DE NS5B (P66) (P70) (RNA-directed RNA polymerase) (EC 2.7.7.48)];
OS Hepatitis C virus (isolate Taiwan) (HCV).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID=31645;
RN [1]
RP SEQUENCE FROM N.A.
RA MEDLINE=9220206; PubMed=1314449;
RA Chen P.J., Lin M.H., Tai K.F., Liu P.C., Lin C.J., Chen D.S.:
RT "The Taiwanese hepatitis C virus genome: sequence determination and
RT mapping the 5' termini of viral genomic and antigenomic RNA.";
RL Virology 189:102-113(1993)
CC -!- FUNCTION: THE SMALL PROTEINS NS2A, NS2B, NS4A AND NS4B ARE
CC HYDROPHOBIC, SUGGESTING A POSSIBLE MEMBRANE-RELATED FUNCTION.
CC NS3 AND NS5 MAY PLAY A ROLE IN THE VIRAL RNA REPLICATION.
CC -!- CATALYTIC ACTIVITY: Hydrolysis of four peptide bonds in the viral
CC precursor polyprotein, commonly with Asp or Glu in the p6
CC position, Cys or Thr in p1 and Ser or Ala in p1'.
CC -!- CATALYTIC ACTIVITY: Nucleoside triphosphate = N diphosphate +
CC [RNA](N).
CC -!- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
CC PROTEIN C AND MRNA.
CC -!- SIMILARITY: THE PROTEASE BELONGS TO PEPTIDASE FAMILY S29.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (see http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: M84754; ; NOT_ANNOTATED_CDS.
CC PIR: A40244; GNMVTV.
CC PDB: 1N64; 25-FEB-98.
CC PDB: 1NS3; 08-APR-98.
CC MEROPS: S29.001; .
CC MEROPS: U39.001; .
CC InterPro: IPR001410; DEAD.
```

```
DR InterPro: IPR002522; HCV_capsid.
DR InterPro: IPR002521; HCV_core.
DR InterPro: IPR002519; HCV_env.
DR InterPro: IPR002531; HCV_NS1.
DR InterPro: IPR002518; HCV_NS2.
DR InterPro: IPR004109; HCV_NS3.
DR InterPro: IPR000745; HCV_NS4a.
DR InterPro: IPR001490; HCV_NS4b.
DR InterPro: IPR002868; HCV_NS5a.
DR InterPro: IPR002166; HCV_RdRP.
DR InterPro: IPR007095; RNA_pol_DS_PS.
DR InterPro: IPR007094; RNA_pol_Psvir.
DR Pfam: PF01543; HCV_core; 1.
DR Pfam: PF01542; HCV_env; 1.
DR Pfam: PF01539; HCV_env; 1.
DR Pfam: PF01560; HCV_NS1; 1.
DR Pfam: PF01538; HCV_NS2; 1.
DR Pfam: PF02907; HCV_NS3; 1.
DR Pfam: PF01006; HCV_NS4a; 1.
DR Pfam: PF01001; HCV_NS4b; 1.
DR Pfam: PF01506; HCV_NS5a; 1.
DR Pfam: PF00271; Helicase_C; 1.
DR Pfam: PF00998; Viral_RdRP; 1.
DR ProDom: PD186062; HCV_NS1; 1.
DR SMARI: SM00487; DEXDC; 1.
DR PolyProtein: Glycoprotein; Transferase; RNA-directed RNA polymerase;
KW Core protein; Coat protein; Envelope protein; Helicase; ATP-binding;
KW Transmembrane; Nonstructural protein; Hydrolase; Serine protease;
KW 3D-structure.
FT INIT_MET 1 1 REMOVED FROM CAPSID PROTEIN C BY THE
FT CHAIN 1 115 CELLULAR AMINOPEPTIDASE..
FT CHAIN 116 191 CORE PROTEIN (POTENTIAL)..
FT CHAIN 192 383 MATRIX PROTEIN (POTENTIAL)..
FT CHAIN 384 729 MAJOR ENVELOPE PROTEIN E (POTENTIAL)..
FT CHAIN 730 1006 NONSTRUCTURAL PROTEIN NS1/E2 (POTENTIAL)..
FT CHAIN 1007 1615 NONSTRUCTURAL PROTEIN NS2 (POTENTIAL)..
FT CHAIN 1616 1862 PROTEASE/HELICASE NS3 (POTENTIAL)..
FT CHAIN 1863 2013 NONSTRUCTURAL PROTEIN NS4A (POTENTIAL)..
FT CHAIN 2014 3010 NONSTRUCTURAL PROTEIN NS4B (POTENTIAL)..
FT TRANSMEM 347 369 RNA-DIRECTED RNA POLYMERASE (POTENTIAL)..
FT ACT_SITE 1083 1087 POTENTIAL.
FT ACT_SITE 1107 1107 CHARGE RELAY SYSTEM (BY SIMILARITY)..
FT ACT_SITE 1165 1165 CHARGE RELAY SYSTEM (BY SIMILARITY)..
FT NP_BIND 1230 1237 ATP (POTENTIAL)..
FT SITE 1316 1319 DECH BOX.
FT CARBOHYD 196 196 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 209 209 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 233 233 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 234 234 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 250 250 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 305 305 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 417 417 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 423 423 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 430 430 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 448 448 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 532 532 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 540 540 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 556 556 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 576 576 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 623 623 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 645 645 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 2041 2041 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 2077 2077 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 2240 2240 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 2529 2529 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 2788 2788 N-LINKED (GLCNAC.. ) (POTENTIAL)..
SQ SEQUENCE 3010 AA; 327047 MW; AAD267D55CDFE215 CRC64;
```

Query Match 93.6%; Score 852; DB 1; Length 3010;
Best Local Similarity 90.1%; Pred. No. 2.7e-77;
Matches 164; Conservative 12; Mismatches 6; Indels 0; Gaps 0;

QY 1 MAPITAYAQOQRLGCGCITTSIGRDKNOVEGEVQIVSTAACTFLAICINGVCWTVYHGA 60
 :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
 Db 1026 LAPITAYAQOQRLGCGCITTSIGRDKNOVEGEVQIVSTAACTFLAICINGVCWTVYHGA 1095
 :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
 QY 51 GERTIASKGPVIQYNTWQDLVWNPAPQGRSLPFCGSSDLYLVTRHADVIVPVRK 120
 :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
 Db 1086 GSKTLAGPKGPTQMYTNDQVLGWHAPQGARSLPFCGSSDLYLVTRHADVIVPVRK 1145
 :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
 QY 121 GSRGSLSPRPISYLKSSGGPLCPACGAVGIFRAAVCGVNAKAVDFIPVESLETTM 180
 :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
 Db 1146 GSRGSLSPRPISYLKSSGGPLCPACGAVGIFRAAVCGVNAKAVDFIPVESLETTM 1205
 :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
 QY 181 RS 192
 :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
 Db 1206 RS 1207
 :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||

RESULT 4

ID POLG_HCVRK STANDARD: PRT: 3010 AA.
 AC P26663;
 DI 01-AUG-1992 (Rel. 23, Created)
 DT 01-AUG-1992 (Rel. 23, Last sequence update)
 DE Genome polyprotein [Contains: Capsid protein; C (Core protein) (#22);
 DE Envelope glycoprotein E1 (GP32) (GP35); Envelope glycoprotein E2
 DE (GP68) (GP70) (NS1); Protein P7; Nonstructural protein NS2 (P21);
 DE (EC 3.4.22.-); Protease/helicase NS3 (P70) (Hepacivirin)
 DE (EC 3.4.21.98); Nonstructural protein NS4A (P4); Nonstructural protein
 DE NS4B (P27); Nonstructural protein NS5A (P56); Nonstructural protein
 DE NS5B (P66) (P70) (RNA-directed RNA polymerase) (EC 2.7.7.48);
 OS Hepatitis C virus (isolate BK) (HCV).
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 OC Hepacivirus;
 OX NCBI_TaxID=11105;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=91140695; PubMed=1847440;
 RA Takamizawa A., Mori C., Fuke I., Manabe S., Murakami S., Fujita J.,
 RA Onishi E., Andoh T., Yoshida I., Okayama H.;
 RT "Structure and organization of the hepatitis C virus genome isolated
 RT from human carriers.";
 RL J. Virol. 65:1205-1213(1991).
 RN [2]
 RP SEQUENCE OF 1487-1500.
 RX MEDLINE=96235224; PubMed=8647104;
 RA Borowski P., Heiland M., Oehmann K., Becker B., Kornetevy L.;
 RT "Non-structural protein 3 of hepatitis C virus inhibits
 RT phosphorylation mediated by CAMP-dependent protein kinase.";
 RL Eur. J. Biochem. 237:611-618(1996).
 RN [3]
 RP X-RAY CRYSTALLOGRAPHY (2.4 ANGSTROMS) OF 1027-1215.
 RX MEDLINE=97015088; PubMed=8861916;
 RA Love R.A., Parage H.E., Wickersham J.A., Hostomsky Z., Habuka N.,
 RA Moenaw E.W., Adachi T., Hostomsky Z.;
 RT "The crystal structure of hepatitis C virus NS3 proteinase reveals a
 RT trypsin-like fold and a structural zinc binding site.";
 RL Cell 87:331-342(1996).
 RN [4]
 RP X-RAY CRYSTALLOGRAPHY (2.2 ANGSTROMS) OF 1027-1210 AND 1578-1631.
 RX MEDLINE=98227846; PubMed=9568891;
 RA Yan Y., Li Y., Mushi S., Sardana V., Cole J.L., Sardana M.,
 RA Steinkuehler C., Tomei L., de Francesco R., Kuo L.C., Chen Z.;
 RT "Complex of NS3 protease and NS4A peptide of BK strain hepatitis C
 RT virus: a 2.2-A resolution structure in a hexagonal crystal form.";
 RL Protein Sci. 7:837-847(1998).
 CC -!- FUNCTION: THE SMALL PROTEINS NS2A, NS2B, NS4A AND NS4B ARE
 CC HYDROPHOBIC, SUGGESTING A POSSIBLE MEMBRANE-RELATED FUNCTION.
 CC NS3 AND NS5 MAY PLAY A ROLE IN THE VIRAL RNA REPLICATION.
 CC -!- CATALYTIC ACTIVITY: Hydrolysis of four peptide bonds in the viral
 CC precursor polyprotein, commonly with Asp or Glu in the P6
 CC position, Cys or Thr in P1 and Ser or Ala in P1'.
 CC -!- CATALYTIC ACTIVITY: N nucleoside triphosphate = N diphosphate +

(RNA)(N).
 -!- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
 CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
 CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
 CC PROTEIN C AND RNA.
 CC -!- SIMILARITY: THE PROTEASE BELONGS TO PEPTIDASE FAMILY S29.
 CC
 CC THIS SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announc/>
 CC or send an email to license@sib-sib.ch).
 CC
 CC EMBL: M58335; AAT72945.1; -
 CC PIR: A38465; GNMVIC.
 CC PDB: 1A10; 25-MAR-98.
 CC PDB: 1JXP; 14-JAN-98.
 CC PDB: 1NS3; 08-APR-98.
 CC PDB: 1C2P; 15-NOV-00.
 CC PDB: 1CS7; 08-NOV-99.
 CC PDB: 1CX5; 09-APR-02.
 CC PDB: 1CX6; 10-APR-02.
 CC PDB: 1QUV; 26-JUN-00.
 CC PDB: 80HM; 20-APR-99.
 CC MEROPS: S29-001; -
 CC MEROPS: U39-061; -
 CC InterPro: IPR001410; DEAD.
 CC InterPro: IPR002522; HCV_capsid.
 CC InterPro: IPR002521; HCV_core.
 CC InterPro: IPR002519; HCV_env.
 CC InterPro: IPR002531; HCV_NS1.
 CC InterPro: IPR002518; HCV_NS2.
 CC InterPro: IPR004109; HCV_NS3.
 CC InterPro: IPR000745; HCV_NS4a.
 CC InterPro: IPR001490; HCV_NS4b.
 CC InterPro: IPR002858; HCV_NS5a.
 CC InterPro: IPR002166; HCV_RdRP.
 CC InterPro: IPR007095; RNA_pol_DS_PS.
 CC InterPro: IPR007094; RNA_pol_PSVir.
 CC Pfam: PF01543; HCV_capsid; 1.
 CC Pfam: PF01542; HCV_core; 1.
 CC Pfam: PF01539; HCV_env; 1.
 CC Pfam: PF01560; HCV_NS1; 1.
 CC Pfam: PF01538; HCV_NS2; 1.
 CC Pfam: PF02907; HCV_NS3; 1.
 CC Pfam: PF01006; HCV_NS4a; 1.
 CC Pfam: PF01001; HCV_NS4b; 1.
 CC Pfam: PF01506; HCV_NS5a; 1.
 CC Pfam: PF00998; Viral_RdRP; 1.
 CC PRODom: PD186062; HCV_NS1; 1.
 CC SMART: SM00487; DEXdc; 1.
 CC Polyprotein; Glycoprotein; Transferase; RNA-directed RNA polymerase;
 CC Core protein; Coat protein; Envelope protein; Helicase; ATP-binding;
 CC Transmembrane; Nonstructural protein; Hydrolase; Serine protease;
 CC 3D-structure.
 CC INIT_MEI 1 ; REMOVED FROM CAPSID PROTEIN C BY THE
 CC CELLULAR AMINOPEPTIDASE.
 CC CHAIN 1 115
 CC CHAIN 116 191
 CC CHAIN 192 383
 CC CHAIN 384 729
 CC CHAIN 730 1006
 CC CHAIN 1007 1615
 CC CHAIN 1616 1862
 CC CHAIN 1863 2013
 CC CHAIN 2014 3010
 CC TRANSMEM 347 369
 CC ACT_SITE 1083 1083
 CC ACT_SITE 1107 1107
 CC ACT_SITE 1165 1165
 CC NP_BIND 1230 1237

FT	1316	1319	DECH BOX.	1316	1319	FT	1316	1319	DECH BOX.
FT	CARBOHYD	196	N-LINKED (GLCNAC)	196	N-LINKED (GLCNAC)	FT	CARBOHYD	196	N-LINKED (GLCNAC)
FT	CARBOHYD	209	N-LINKED (GLCNAC)	209	N-LINKED (GLCNAC)	FT	CARBOHYD	209	N-LINKED (GLCNAC)
FT	CARBOHYD	234	N-LINKED (GLCNAC)	234	N-LINKED (GLCNAC)	FT	CARBOHYD	234	N-LINKED (GLCNAC)
FT	CARBOHYD	250	N-LINKED (GLCNAC)	250	N-LINKED (GLCNAC)	FT	CARBOHYD	250	N-LINKED (GLCNAC)
FT	CARBOHYD	305	N-LINKED (GLCNAC)	305	N-LINKED (GLCNAC)	FT	CARBOHYD	305	N-LINKED (GLCNAC)
FT	CARBOHYD	417	N-LINKED (GLCNAC)	417	N-LINKED (GLCNAC)	FT	CARBOHYD	417	N-LINKED (GLCNAC)
FT	CARBOHYD	423	N-LINKED (GLCNAC)	423	N-LINKED (GLCNAC)	FT	CARBOHYD	423	N-LINKED (GLCNAC)
FT	CARBOHYD	430	N-LINKED (GLCNAC)	430	N-LINKED (GLCNAC)	FT	CARBOHYD	430	N-LINKED (GLCNAC)
FT	CARBOHYD	448	N-LINKED (GLCNAC)	448	N-LINKED (GLCNAC)	FT	CARBOHYD	448	N-LINKED (GLCNAC)
FT	CARBOHYD	532	N-LINKED (GLCNAC)	532	N-LINKED (GLCNAC)	FT	CARBOHYD	532	N-LINKED (GLCNAC)
FT	CARBOHYD	540	N-LINKED (GLCNAC)	540	N-LINKED (GLCNAC)	FT	CARBOHYD	540	N-LINKED (GLCNAC)
FT	CARBOHYD	556	N-LINKED (GLCNAC)	556	N-LINKED (GLCNAC)	FT	CARBOHYD	556	N-LINKED (GLCNAC)
FT	CARBOHYD	576	N-LINKED (GLCNAC)	576	N-LINKED (GLCNAC)	FT	CARBOHYD	576	N-LINKED (GLCNAC)
FT	CARBOHYD	623	N-LINKED (GLCNAC)	623	N-LINKED (GLCNAC)	FT	CARBOHYD	623	N-LINKED (GLCNAC)
FT	CARBOHYD	645	N-LINKED (GLCNAC)	645	N-LINKED (GLCNAC)	FT	CARBOHYD	645	N-LINKED (GLCNAC)
FT	CARBOHYD	2041	N-LINKED (GLCNAC)	2041	N-LINKED (GLCNAC)	FT	CARBOHYD	2041	N-LINKED (GLCNAC)
FT	CARBOHYD	2077	N-LINKED (GLCNAC)	2077	N-LINKED (GLCNAC)	FT	CARBOHYD	2077	N-LINKED (GLCNAC)
FT	CARBOHYD	2240	N-LINKED (GLCNAC)	2240	N-LINKED (GLCNAC)	FT	CARBOHYD	2240	N-LINKED (GLCNAC)
FT	CARBOHYD	2529	N-LINKED (GLCNAC)	2529	N-LINKED (GLCNAC)	FT	CARBOHYD	2529	N-LINKED (GLCNAC)
FT	CARBOHYD	2788	N-LINKED (GLCNAC)	2788	N-LINKED (GLCNAC)	FT	CARBOHYD	2788	N-LINKED (GLCNAC)
FT	STRAND	1031	N-LINKED (GLCNAC)	1031	N-LINKED (GLCNAC)	FT	STRAND	1031	N-LINKED (GLCNAC)
FT	HELIX	1039	N-LINKED (GLCNAC)	1039	N-LINKED (GLCNAC)	FT	HELIX	1039	N-LINKED (GLCNAC)
FT	STRAND	1050	N-LINKED (GLCNAC)	1050	N-LINKED (GLCNAC)	FT	STRAND	1050	N-LINKED (GLCNAC)
FT	STRAND	1059	N-LINKED (GLCNAC)	1059	N-LINKED (GLCNAC)	FT	STRAND	1059	N-LINKED (GLCNAC)
FT	TURN	1068	N-LINKED (GLCNAC)	1068	N-LINKED (GLCNAC)	FT	TURN	1068	N-LINKED (GLCNAC)
FT	TURN	1075	N-LINKED (GLCNAC)	1075	N-LINKED (GLCNAC)	FT	TURN	1075	N-LINKED (GLCNAC)
FT	STRAND	1077	N-LINKED (GLCNAC)	1077	N-LINKED (GLCNAC)	FT	STRAND	1077	N-LINKED (GLCNAC)
FT	HELIX	1082	N-LINKED (GLCNAC)	1082	N-LINKED (GLCNAC)	FT	HELIX	1082	N-LINKED (GLCNAC)
FT	TURN	1086	N-LINKED (GLCNAC)	1086	N-LINKED (GLCNAC)	FT	TURN	1086	N-LINKED (GLCNAC)
FT	TURN	1090	N-LINKED (GLCNAC)	1090	N-LINKED (GLCNAC)	FT	TURN	1090	N-LINKED (GLCNAC)
FT	STRAND	1093	N-LINKED (GLCNAC)	1093	N-LINKED (GLCNAC)	FT	STRAND	1093	N-LINKED (GLCNAC)
FT	STRAND	1095	N-LINKED (GLCNAC)	1095	N-LINKED (GLCNAC)	FT	STRAND	1095	N-LINKED (GLCNAC)
FT	TURN	1101	N-LINKED (GLCNAC)	1101	N-LINKED (GLCNAC)	FT	TURN	1101	N-LINKED (GLCNAC)
FT	TURN	1104	N-LINKED (GLCNAC)	1104	N-LINKED (GLCNAC)	FT	TURN	1104	N-LINKED (GLCNAC)
FT	STRAND	1108	N-LINKED (GLCNAC)	1108	N-LINKED (GLCNAC)	FT	STRAND	1108	N-LINKED (GLCNAC)
FT	STRAND	1120	N-LINKED (GLCNAC)	1120	N-LINKED (GLCNAC)	FT	STRAND	1120	N-LINKED (GLCNAC)
FT	STRAND	1122	N-LINKED (GLCNAC)	1122	N-LINKED (GLCNAC)	FT	STRAND	1122	N-LINKED (GLCNAC)
FT	TURN	1129	N-LINKED (GLCNAC)	1129	N-LINKED (GLCNAC)	FT	TURN	1129	N-LINKED (GLCNAC)
FT	TURN	1135	N-LINKED (GLCNAC)	1135	N-LINKED (GLCNAC)	FT	TURN	1135	N-LINKED (GLCNAC)
FT	STRAND	1139	N-LINKED (GLCNAC)	1139	N-LINKED (GLCNAC)	FT	STRAND	1139	N-LINKED (GLCNAC)
FT	STRAND	1149	N-LINKED (GLCNAC)	1149	N-LINKED (GLCNAC)	FT	STRAND	1149	N-LINKED (GLCNAC)
FT	HELIX	1158	N-LINKED (GLCNAC)	1158	N-LINKED (GLCNAC)	FT	HELIX	1158	N-LINKED (GLCNAC)
FT	TURN	1162	N-LINKED (GLCNAC)	1162	N-LINKED (GLCNAC)	FT	TURN	1162	N-LINKED (GLCNAC)
FT	TURN	1165	N-LINKED (GLCNAC)	1165	N-LINKED (GLCNAC)	FT	TURN	1165	N-LINKED (GLCNAC)
FT	STRAND	1168	N-LINKED (GLCNAC)	1168	N-LINKED (GLCNAC)	FT	STRAND	1168	N-LINKED (GLCNAC)
FT	TURN	1172	N-LINKED (GLCNAC)	1172	N-LINKED (GLCNAC)	FT	TURN	1172	N-LINKED (GLCNAC)
FT	STRAND	1175	N-LINKED (GLCNAC)	1175	N-LINKED (GLCNAC)	FT	STRAND	1175	N-LINKED (GLCNAC)
FT	TURN	1187	N-LINKED (GLCNAC)	1187	N-LINKED (GLCNAC)	FT	TURN	1187	N-LINKED (GLCNAC)
FT	STRAND	1189	N-LINKED (GLCNAC)	1189	N-LINKED (GLCNAC)	FT	STRAND	1189	N-LINKED (GLCNAC)
FT	HELIX	1198	N-LINKED (GLCNAC)	1198	N-LINKED (GLCNAC)	FT	HELIX	1198	N-LINKED (GLCNAC)
FT	TURN	1203	N-LINKED (GLCNAC)	1203	N-LINKED (GLCNAC)	FT	TURN	1203	N-LINKED (GLCNAC)
FT	STRAND	1680	N-LINKED (GLCNAC)	1680	N-LINKED (GLCNAC)	FT	STRAND	1680	N-LINKED (GLCNAC)
SQ	SEQUENCE	3010 AA		327165 MW	F8422D5ECCDFD9C3RC64;	SQ	SEQUENCE	3010 AA	
Query Match 93.5% Score 991; DB 1; length 3010;									
Rest Local Similarity 89.0%; Pred. No. 3.4e-77;									
Matches 162; Conservative 15; Mismatches 5; Gaps 0; Gaps 0;									
Qy	1	MALITAYAOQTRG:LGCIITSLTRCKNQVEGVQVSVIAACFLATCINGVCWTVYHGA	50	1	MALITAYAOQTRG:LGCIITSLTRCKNQVEGVQVSVIAACFLATCINGVCWTVYHGA	Qy	1	MALITAYAOQTRG:LGCIITSLTRCKNQVEGVQVSVIAACFLATCINGVCWTVYHGA	50
Db	1026	LAPITAYSOQTRG:LGCIITSLTRCKNQVEGVQVSVIAACFLATCINGVCWTVYHGA	1085	1026	LAPITAYSOQTRG:LGCIITSLTRCKNQVEGVQVSVIAACFLATCINGVCWTVYHGA	Db	1026	LAPITAYSOQTRG:LGCIITSLTRCKNQVEGVQVSVIAACFLATCINGVCWTVYHGA	1085
Qy	61	GTRITASPKGFVIMTVNVDKLVGWPAPOGSRSLTPTCTGSSDLVLTTRHADVIPVRRR	120	61	GTRITASPKGFVIMTVNVDKLVGWPAPOGSRSLTPTCTGSSDLVLTTRHADVIPVRRR	Qy	61	GTRITASPKGFVIMTVNVDKLVGWPAPOGSRSLTPTCTGSSDLVLTTRHADVIPVRRR	120
Db	1086	GSKTLAAPKGPITOMY:NVQDLVGVWPKPGCARSLTPTCTGSSDLVLTTRHADVIPVRRR	1145	1086	GSKTLAAPKGPITOMY:NVQDLVGVWPKPGCARSLTPTCTGSSDLVLTTRHADVIPVRRR	Db	1086	GSKTLAAPKGPITOMY:NVQDLVGVWPKPGCARSLTPTCTGSSDLVLTTRHADVIPVRRR	1145
Qy	12	GDSRGSLLSPRPISY:JAGSGGGPLLCVAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM	180	12	GDSRGSLLSPRPISY:JAGSGGGPLLCVAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM	Qy	12	GDSRGSLLSPRPISY:JAGSGGGPLLCVAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM	180
Db	1146	GDSRGSLLSPRPISY:JAGSGGGPLLCVAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM	1205	1146	GDSRGSLLSPRPISY:JAGSGGGPLLCVAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM	Db	1146	GDSRGSLLSPRPISY:JAGSGGGPLLCVAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM	1205
Qy	181	RS 182		181	RS 182	Qy	181	RS 182	

UB	1206	RS 1207	UB	1206	RS 1207
RESULT 5			RESULT 5		
POLG_HCV7A			POLG_HCV7A		
CD	POLG_HCV7A	STANDARD; PRT: 3010 AA.	CD	POLG_HCV7A	STANDARD; PRT: 3010 AA.
AC	P26662;		AC	P26662;	
DT	01-AUG-1992 (Rel. 23, Created)		DT	01-AUG-1992 (Rel. 23, Created)	
DT	01-AUG-1992 (Rel. 23, Last sequence update)		DT	01-AUG-1992 (Rel. 23, Last sequence update)	
DE	Genome polyprotein [Contains: Capsid protein C (Core protein) (P22); Envelope glycoprotein E1 (GP32) (GP35); Envelope glycoprotein E2 (GP68) (GP70) (NS1); Protein P7; Nonstructural protein NS2 (P21) (EC 3.4.22.-); Protease/helicase NS3 (P70) (Hepacivirin) (EC 3.4.22.-); Nonstructural protein NS4A (P4); Nonstructural protein NS4B (P27); Nonstructural protein NS5A (P56); Nonstructural protein NS5B (P66) (P70) (RNA-directed RNA polymerase) (HCV).]		DE	Genome polyprotein [Contains: Capsid protein C (Core protein) (P22); Envelope glycoprotein E1 (GP32) (GP35); Envelope glycoprotein E2 (GP68) (GP70) (NS1); Protein P7; Nonstructural protein NS2 (P21) (EC 3.4.22.-); Protease/helicase NS3 (P70) (Hepacivirin) (EC 3.4.22.-); Nonstructural protein NS4A (P4); Nonstructural protein NS4B (P27); Nonstructural protein NS5A (P56); Nonstructural protein NS5B (P66) (P70) (RNA-directed RNA polymerase) (HCV).]	
OS	Hepatitis C virus (isolate Japanese) (HCV).		OS	Hepatitis C virus (isolate Japanese) (HCV).	
OC	Viruses; ssRNA positive-strand viruses, no DNA stage: Flaviviridae; Hepacivirus.		OC	Viruses; ssRNA positive-strand viruses, no DNA stage: Flaviviridae; Hepacivirus.	
OX	NCBI_TaxID=11116;		OX	NCBI_TaxID=11116;	
RN	[1]		RN	[1]	
RP	SEQUENCE FROM N.A.		RP	SEQUENCE FROM N.A.	
RX	MEDLINE=9108550; PubMed=2175903;		RX	MEDLINE=9108550; PubMed=2175903;	
RA	Kato N., Hijikata M., Ootsuyama Y., Nakagawa M., Ohkoshi S., Sugimura T., Shimotohno K.;		RA	Kato N., Hijikata M., Ootsuyama Y., Nakagawa M., Ohkoshi S., Sugimura T., Shimotohno K.;	
RT	"Molecular cloning of the human hepatitis C virus genome from Japanese patients with non-A, non-B hepatitis.";		RT	"Molecular cloning of the human hepatitis C virus genome from Japanese patients with non-A, non-B hepatitis.";	
RL	Proc. Natl. Acad. Sci. U.S.A. 87:9524-9528(1990).		RL	Proc. Natl. Acad. Sci. U.S.A. 87:9524-9528(1990).	
RN	[2]		RN	[2]	
RP	DISCUSSION OF SEQUENCE.		RP	DISCUSSION OF SEQUENCE.	
RX	MEDLINE=91192160; PubMed=1849488;		RX	MEDLINE=91192160; PubMed=1849488;	
RA	Kato N., Hijikata M., Nakagawa M., Ootsuyama Y., Muraiso K., Ohkoshi S., Shimotohno K.;		RA	Kato N., Hijikata M., Nakagawa M., Ootsuyama Y., Muraiso K., Ohkoshi S., Shimotohno K.;	
RT	"Molecular structure of the Japanese hepatitis C viral genome.";		RT	"Molecular structure of the Japanese hepatitis C viral genome.";	
CC	FEBS Lett. 280:325-328(1991).		CC	FEBS Lett. 280:325-328(1991).	
CC	FUNCTION: THE SMALL PROTEINS NS2A, NS2B, NS4A AND NS4B ARE HYDROPHOBIC, SUGGESTING A POSSIBLE MEMBRANE-RELATED FUNCTION.		CC	FUNCTION: THE SMALL PROTEINS NS2A, NS2B, NS4A AND NS4B ARE HYDROPHOBIC, SUGGESTING A POSSIBLE MEMBRANE-RELATED FUNCTION.	
CC	NS3 AND NS5 MAY PLAY A ROLE IN THE VIRAL RNA REPLICATION.		CC	NS3 AND NS5 MAY PLAY A ROLE IN THE VIRAL RNA REPLICATION.	
CC	!- CATALYTIC ACTIVITY: Hydrolysis of four peptide bonds in the viral precursor polyprotein, commonly with Asp or Glu in the P6 position, Cys or Thr in P1 and Ser or Ala in P1'.		CC	!- CATALYTIC ACTIVITY: Hydrolysis of four peptide bonds in the viral precursor polyprotein, commonly with Asp or Glu in the P6 position, Cys or Thr in P1 and Ser or Ala in P1'.	
CC	!- CATALYTIC ACTIVITY: N nucleoside triphosphate - N diphosphate + {RNA}(N).		CC	!- CATALYTIC ACTIVITY: N nucleoside triphosphate - N diphosphate + {RNA}(N).	
CC	!- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS: PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF PROTEIN C AND MRNA.		CC	!- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS: PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF PROTEIN C AND MRNA.	
CC	!- SIMILARITY: THE PROTEASE BELONGS TO PEPTIDASE FAMILY S29.		CC	!- SIMILARITY: THE PROTEASE BELONGS TO PEPTIDASE FAMILY S29.	
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See http://www.isb-sib.ch/annouce/ or send an email to licenses@isb-sib.ch).		CC	This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See http://www.isb-sib.ch/annouce/ or send an email to licenses@isb-sib.ch).	
CC	EMBL: D90208; BAAL4233.1; -		CC	EMBL: D90208; BAAL4233.1; -	
DR	PIR: A35253; GNWVCL.		DR	PIR: A35253; GNWVCL.	
DR	HSP: P26663; LUXP.		DR	HSP: P26663; LUXP.	
DR	MEROPS: S29.001; -		DR	MEROPS: S29.001; -	
DR	MEROPS: U39.001; -		DR	MEROPS: U39.001; -	
DR	InterPro: IPR001410; DEAD.		DR	InterPro: IPR001410; DEAD.	
DR	InterPro: IPR002522; HCV_capsid.		DR	InterPro: IPR002522; HCV_capsid.	
DR	InterPro: IPR002521; HCV_core.		DR	InterPro: IPR002521; HCV_core.	
DR	InterPro: IPR002519; HCV_env.		DR	InterPro: IPR002519; HCV_env.	
DR	InterPro: IPR002531; HCV_NSI.		DR	InterPro: IPR002531; HCV_NSI.	
DR	InterPro: IPR002518; HCV_NS2.		DR	InterPro: IPR002518; HCV_NS2.	
DR	InterPro: IPR004109; HCV_NS3.		DR	InterPro: IPR004109; HCV_NS3.	
DR	InterPro: IPR000745; HCV_NS4a.		DR	InterPro: IPR000745; HCV_NS4a.	
DR	InterPro: IPR001490; HCV_NS4b.		DR	InterPro: IPR001490; HCV_NS4b.	
DR	InterPro: IPR002868; HCV_NS5a.		DR	InterPro: IPR002868; HCV_NS5a.	

DR	Pfam:	PF01543:	HCV_capsid; 1.
DR	Pfam:	PF01542:	HCV_core; 1.
DR	Pfam:	PF01539:	HCV_env; 1.
DR	Pfam:	PF01560:	HCV_NS1; 1.
DR	Pfam:	PF01538:	HCV_NS2; 1.
DR	Pfam:	PF02907:	HCV_NS3; 1.
DR	Pfam:	PF01006:	HCV_NS4a; 1.
DR	Pfam:	PF01001:	HCV_NS4b; 1.
DR	Pfam:	PF01506:	HCV_NS5a; 1.
DR	Pfam:	PF00271:	helicase-C; 1.
DR	Pfam:	PF00998:	Viral_RdRP; 1.
DR	Pfam:	PF0186062:	HCV_NS1; 1.
DR	SMART:	SM03487:	DRDC; 1.
KW	Polyprotein:		Transferase: RNA-directed RNA polymerase;
KW	Core protein:		Envelope protein: Helicase: ATP-binding;
KW	Transmembrane:		Nonstructural protein; Hydrolase: Serine protease;
KW	3D-structure:		
FT	INIT_MET	1	REMOVED FROM CAPSID PROTEIN C BY THE CELLULAR AMINOPEPTIDASE.
FT	CHAIN	1	CAPSID PROTEIN C (POTENTIAL).
FT	CHAIN	116	MATRIX PROTEIN (POTENTIAL).
FT	CHAIN	192	MAJOR ENVELOPE PROTEIN E (POTENTIAL).
FT	CHAIN	384	NONSTRUCTURAL PROTEIN NS1/E2 (POTENTIAL).
FT	CHAIN	730	NON-STRUCTURAL PROTEIN NS2 (POTENTIAL).
FT	CHAIN	1007	PROTEASE/HELICASE NS3 (POTENTIAL).
FT	CHAIN	1616	NONSTRUCTURAL PROTEIN NS4A (POTENTIAL).
FT	CHAIN	1863	NONSTRUCTURAL PROTEIN NS4B (POTENTIAL).
FT	CHAIN	2014	RNA-DIRECTED RNA POLYMERASE (POTENTIAL).
FT	CHAIN	347	POTENTIAL.
FT	TRANSMEM	1083	CHARGE RELAY SYSTEM (BY SIMILARITY).
FT	ACT_SITE	1107	CHARGE RELAY SYSTEM (BY SIMILARITY).
FT	ACT_SITE	1230	CHARGE RELAY SYSTEM (BY SIMILARITY).
FT	NP_BIND	1230	ATP (POTENTIAL).
FT	SITE	1316	DECH BOX.
FT	CARBOHYD	196	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	209	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	234	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	250	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	305	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	417	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	423	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	430	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	448	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	532	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	540	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	556	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	576	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	623	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	645	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	2041	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	2077	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	2240	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	2529	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	2788	N-LINKED (GLCNAC) (POTENTIAL).
FT	CARBOHYD	3010	N-LINKED (GLCNAC) (POTENTIAL).
SQ	SEQUENCE	3010 AA:	325573 MW; 94A1C77435D642B5 CRC64;

Query Match	92.88:	Score 384:	DB 1:	Length 3010:
Best Local Similarity	88.56:	Pred. No. 1.5e-76:		
Matches 161:	Conservative 15:	Mismatches 5:	Indels 0:	Gaps 0:
QY	1	MAPTAAQOTRGLGCIITSLTGROKNQVEGVQVSVTAACTEFLATPCINGVQWTVHGA	60	
DB	1026	LAPITAAQOTRGLGCIIVISLTGRUKNQVEGVQVSVTAQSFATPCVNGVQWTVHGA	1065	
QY	61	GTRTASPKGVQIOMYTNVQKDLGVMPAPQGGSRSLTPCTCGSSDLXLVTRHADVIPVRRR	120	
DB	1086	GSKILAGEKGPITOMYTNVQDLGVMPAPQGGSRSLTPCTCGSSDLXLVTRHADVIPVRRR	1145	
QY	121	GDGRGSLSPRPISYKCGSSGGLLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLTTM	180	
DB	1146	GDGRGSLSPRPVSYKCGSSGGLLCPGSHAGVIFRAAVCTRGVAKAVDFIPVESMETTM	1205	
QY	181	RS	182	

DB	1206	RS	1207
POLG_HCVJ8		STANDARD:	PRT: 3033 AA.
ID	P26661:		
AC	01-AUG-1992 (Rel. 23, Created)		
DT	01-AUG-1992 (Rel. 23, Last sequence update)		
DT	28-FEB-2003 (Rel. 41, Last annotation update)		
DE	Genome polyprotein [Contains: Capsid protein C (Core protein) (P22); Envelope glycoprotein E1 (GP32); Envelope glycoprotein E2 (GP58) (GP70) (NS1); Protein P7; Nonstructural protein NS2 (P21) (EC 3.4.22.-); Protease/helicase NS3 (P70) (Hepacivirin) (EC 3.4.21.98); Nonstructural protein NS4A (P4); Nonstructural protein NS4B (P27); Nonstructural protein NS5A (P56); Nonstructural protein NS5B (P66) (P70) (RNA-directed RNA polymerase) (EC 2.7.7.48)].		
OS	Hepatitis C virus (isolate HC-J8) (HCV).		
OC	Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae; Hepacivirus.		
OX	NCBI_TaxID=11115;		
RN	[1]_TaxID=11115;		
RP	SEQUENCE FROM N.A.		
RX	MEDLINE=92230322; PubMed=1314459;		
RA	Okamoto H., Kurai K., Okada S.-I., Yamamoto K., Lizuka H., Tanaka T., Fukuda S., Tsuda F., Mishiro S.;		
RT	*Full-length sequence of a hepatitis C virus genome having poor homology to reported isolates: comparative study of four distinct genotypes.*;		
RT	Virology 188:331-341(1992).		
KL	FUNCTION: THE SMALL PROTEINS NS2A, NS2B, NS4A AND NS4B ARE HYDROPHOBIC, SUGGESTING A POSSIBLE MEMBRANE-RELATED FUNCTION.		
CC	NS3 AND NS5 MAY PLAY A ROLE IN THE VIRAL RNA REPLICATION.		
CC	!- CATALYTIC ACTIVITY: Hydrolysis of four peptide bonds in the viral precursor polyprotein, commonly with Asp or Glu in the P6 position. Cys or Thr in P1 and Ser or Ala in P1'.		
CC	!- CATALYTIC ACTIVITY: N nucleoside triphosphate = N diphosphate + [RNA](N).		
CC	!- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS: PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF PROTEIN C AND MRNA.		
CC	!- SIMILARITY: THE PROTEASE BELONGS TO PEPTIDASE FAMILY S29.		
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL Outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See http://www.isb-sib.ch/announce/ or send an email to license@sib-sib.ch).		
DR	EMBL: J10988; BAA01761.1;		
DR	PIR: A40250; GNMVJ8.		
DR	HSSP: P27958; IHEI.		
DR	MEROPS: S29.001; -.		
DR	MEROPS: U39.001; -.		
DR	InterPro: IPR001410; DEAD.		
DR	InterPro: IPR002522; HCV_capsid.		
DR	InterPro: IPR002521; HCV_core.		
DR	InterPro: IPR002519; HCV_env.		
DR	InterPro: IPR002531; HCV_NS1.		
DR	InterPro: IPR002518; HCV_NS2.		
DR	InterPro: IPR004109; HCV_NS3.		
DR	InterPro: IPR000745; HCV_NS4a.		
DR	InterPro: IPR001490; HCV_NS4b.		
DR	InterPro: IPR002868; HCV_NS5a.		
DR	InterPro: IPR002166; HCV_RdRP.		
DR	InterPro: IPR007095; RNA_pol_DS_PS.		
DR	InterPro: IPR007094; RNA_pol_PSVir.		
DR	Pfam: PF01543; HCV_capsid; 1.		
DR	Pfam: PF01542; HCV_core; 1.		

Pfam: PF02907; HCV_NS3; 1.
 Pfam: PF01006; HCV_NS4a; 1.
 Pfam: PF01901; HCV_NS4b; 1.
 Pfam: PF01536; HCV_NS5a; 1.
 Pfam: PF00271; helicase_C; 1.
 Pfam: PF00998; Viral_RDRP; 1.
 ProDom: PD185062; HCV_NS1; 1.
 SMART: SM00487; DEXDC; 1.
 Polyprotein; Glycoprotein; Transferase; RNA-directed RNA polymerase;
 Core protein; Coat protein; Envelope protein; Helicase; ATP-binding;
 Transmembrane; Nonstructural proteins; Hydrolase; Serine protease.
 INIT_MET 1
 FT CHAIN 1 125
 FT CHAIN 191
 FT CHAIN 192 393
 FT CHAIN 384 733
 FT CHAIN 734 1010
 FT CHAIN 1011 1619
 FT CHAIN 1620 1866
 FT CHAIN 1867 2017
 FT CHAIN 2018 3033
 FT TRANSMEM 347 369
 FT ACT_SITE 1087 1087
 FT ACT_SITE 1111 1111
 FT ACT_SITE 1169 1169
 FT NP_BIND 1234 1241
 FT SITE 1320 1323
 FT CARBOHYD 196 196
 FT CARBOHYD 209 209
 FT CARBOHYD 234 234
 FT CARBOHYD 305 305
 FT CARBOHYD 417 417
 FT CARBOHYD 423 423
 FT CARBOHYD 436 436
 FT CARBOHYD 448 448
 FT CARBOHYD 477 477
 FT CARBOHYD 534 534
 FT CARBOHYD 542 542
 FT CARBOHYD 558 558
 FT CARBOHYD 578 578
 FT CARBOHYD 627 627
 FT CARBOHYD 649 649
 FT CARBOHYD 1091 1091
 FT CARBOHYD 2038 2038
 FT CARBOHYD 2811 2811
 SQ SEQUENCE 3033 AA; 329155 MW; F957F5C1A273BE9F CAC64;
 Query Match 74.74; Score 712; DR 1; Length 3033;
 Best Local Similarity 69.88; Pred. No. 5.3e-60;
 Matches 127; Conservative 29; Mismatches 26; Indels 0; Gaps 0;
 QY 1 MAPITAYAAQTGRLGCTTSLTGRDNQVEGFVOIVSTAAQTFLATPCINGVWTVYHGA 60
 Db 1030 LAPTAYAAQTGRLGCTTSLTGRDNQVEGFVOIVSTAAQTFLATPCINGVWTVYHGA 1069
 QY 61 GTRTIASPKGPVQIOMYTNVTKDLVGNPAPQGSRLTPCTCGSSDLYLVTHAVIPVRRR 120
 Db 1090 GNKTLAGSRGPVQIOMYSSAEGDLVGNPAPQGSRLTPCTCGSSDLYLVTHAVIPVRRR 1149
 QY 121 GDSRGSLLSPRISLYLKGSSGGPGLLCGAGVAGVIFRAAVCTRGVAKAVDFIPVESLET 150
 Db 1150 GDRKCALLSPRLSTLKGSSGGPGLLCGAGVAGVIFRAAVCTRGVAKAVDFIPVESLET 1209
 QY 181 RS 182
 Db 1210 RS 1211
 RESULT 9
 ID PAAD_PSEAE
 AC C9HX08; STANDARD; PRI: 209 AA.

DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Probable aromatic acid decarboxylase (EC 4.1.1.1).
 GN PA4019.
 OS Pseudomonas aeruginosa.
 CC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
 CC Pseudomonadaceae; Pseudomonas.
 CX NCBI_TaxID=287;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=ATCC 15692 / PAO.;
 RX MEDLINE=20437337; PubMed=10984043;
 RA Stover C.K., Pham X.-Q.T., Erwin A.L., Mizoguchi S.D., Warren P.,
 RA Hickey M.J., Brinkman F.S.L., Hufnagle W.O., Kowalik D.J., Lagrou M.,
 RA Garber R.J., Goltz L., Tolentino E., Westbrock-Wadman S., Yuan Y.,
 RA Brody L.L., Coulter S.N., Folger K.R., Kas A., Larbig K., Lim R.M.,
 RA Smith K.A., Spencer D.H., Wong G.K.-S., Wu Z., Paulsen I.T.,
 RA Reizer J., Sailer M.H., Hancock R.E.W., Lory S., Olson M.V.;
 RI "Complete genome sequence of Pseudomonas aeruginosa PAO1, an
 RI opportunistic pathogen";
 RL Nature 406:959-964 (2000).
 CC -1- SIMILARITY: BELONGS TO THE POLYPRENYL P-HYDROXYBENZOATE /
 CC PHENYLACRYLIC ACID DECARBOXYLASES FAMILY.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement. (See http://www.isb-sib.ch/announce/
 CC or send an email to license@sib-sib.ch).
 CC -----
 CC EMBL: AF004618; AAG07406.1; .
 DR PIR: H83144; H83144.
 DR InterPro: IPR003382; Flavoprotein.
 DR Pfam: PF02441; Flavoprotein: 1.
 KW Hypothetical protein; Lyase; Decarboxylase; Complete proteome.
 SQ SEQUENCE 209 AA; 22367 MW; 01FD081CC495D3F6 CRC64;
 Query Match 9.11; Score 87; DR 1; Length 209;
 Best Local Similarity 26.28; Pred. No. 0.28;
 Matches 55; Conservative 18; Mismatches 61; Indels 76; Gaps 12;
 QY 8 AQQTGRLGCTTSLTGRDNQVEGFVOIVSTAAQTFLATPCINGVWTVYHGA 66
 Db 17 AQVGLRLDCLV-----QFEVHPLISKAAQLVMAI-----ETVVA 53
 QY 67 SPKGP-----VIQMTNVKDLVGNPAPQGSRLTP-----CTCGSSDL 105
 Db 54 LPAKFOAQMAQFLTEXGAAAGQIRVFGQND-----WNAPPASGSGAPNAWICPCSTGTL 108
 QY 106 -----YLVRGADVTPVRRGDSRGSLLSPR--PTS-----YLKSGSGPPLCPA 148
 Db 109 SAVATGACACNLLERRADVALKER---RPLVLVPREAFSSIHLENMILKSLNLGAVILPA 164
 QY 149 GHAVGIFRAAVCTRGVAKAVDFIPVESLET 178
 Db 165 --APGFYHQ---PQSVEDLDLVFVVAAILNT 189
 RESULT 10
 ID HHOA_ARATH
 AC Q9SEL7; O49507; STANDARD; PRI: 321 AA.
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE protease HhoA, chloroplast precursor (EC 3.4.21.-).
 GN HHOA OR AT4G18370 OR F28J12.30.
 OS Arabidopsis thaliana (Mouse-ear cress).
 CC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 CC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicotyledons; Rosidae;

CC eutrods II; Brassicales; Brassicaceae; Arabidopsis.
 CC NCBI_TaxID=3702;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Lersch M.H.A., Sokolenko A., Herrmann R.G.:
 RT "Identification and characterization of the chloroplast HspA protease,
 RT a homolog to the bacterial periplasmic protease HspA";
 RL submitted (DEC-1998) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=cv. Columbia;
 RX MEDLINE=26083488; PubMed=10617198;
 RA Mayer K.F.X., Schueller C., Wambutt R., Murphy G., Volckaert G.,
 RA Pohl T., Duesterhoeft A., Stiekema W., Estian K.-D., Terryn N.,
 RA Harris B., Anstoe W., Brandt P., Grivell L., Kieger M.,
 RA Weichselgartner M., de Simone V., Obermaier B., Maché R., Mueller M.,
 RA Kreis M., Delsen M., Pulgomech P., Watson M., Schmidkeisen I.,
 RA Reichert B., Portetle E., Perez-Alonso M., Souty M., Bancroft T.,
 RA Vos P., Honeisel C., Zimmermann W., Wadler H., Fildes P.,
 RA Langham S.-A., McCullagh B., Blum L., Robben J.,
 RA Van der Schueren J., Grymoprez B., Chuang Y.-J., Vandenbussche P.,
 RA Bracken M., Weltyons J., Voet M., Bastiaens J., Aert R., DeCoor E.,
 RA Weitzenecker T., Bothe G., Ransperger J., Hilbert H., Braun M.,
 RA Holzer E., Brandt A., Peters S., van Staveren M., Dirkse W.,
 RA Moolman P., Klein Lankhorst R., Rose M., Hauf J., Koetler P.,
 RA Berner S., Hempel S., Fedpausch M., Lambert S., Van den Daele H.,
 RA De Keyser A., Huysschaert C., Giesch J., Villarroel R., De Clercq R.,
 RA Van Montagu M., Kogers C., Cronin A., Qail M., Bray-Allen S.,
 RA Clark L., Dorgett J., Hall S., Kay M., Leonard N., McLay K., Mayes R.,
 RA Pettelt A., Rajandream M.A., Lyne M., Boncs V., Rechmann S.,
 RA Borkova D., Bloeker A., Scharie M., Grimm M., Koehnert T.-H.,
 RA Dose S., de Haan M., Maarse A., Schaefer M., Mueller-Auer S.,
 RA Gabel C., Fuchs M., Fartmann B., Grandrath K., Dauner D., Herzi A.,
 RA Neumann S., Argiriou A., Vitale D., Liguori R., Piravandi E.,
 RA Massenot O., Quigley F., Clabaud G., Kuendlein A., Felber R.,
 RA Schnabel F., Hiller R., Schmidt W., Lecharny A., Abouit S.,
 RA Chedrol F., Cooke R., Berger C., Monfort A., Casacuberta E.,
 RA Gibbons T., Weber N., Vandenbol M., Barques M., Terol C., Iorres A.,
 RA Perez-Perez A., Purnelle B., Best E., Johnson S., Tabor D., Jesse T.,
 RA Heijnen L., Schwarz S., Scholler P., Heber S., Francis P., Bieleke C.,
 RA Frishman D., Haase D., Lemcke K., Meves H.-W., Stocker S.,
 RA Zaccaria P., Bevan M., Wilson R.K., de la Bastide M., Habermann K.,
 RA Parnell L., Dedhia N., Grol L., Schutz K., Huang E., Spiegel L.,
 RA Sekhon M., Murray J., Sheet P., Cordes M., Abu-Threidh J.,
 RA Stoneking T., Kalicki J., Graves T., Harmon G., Edwards J.,
 RA Latreille P., Courtney L., Cloud J., Abbott A., Scott K., Johnson D.,
 RA Minx P., Bentley D., Fulton B., Miller N., Greco T., Kemp K.,
 RA Kramer J., Fulton L., Mardis E., Dante M., Pepin K., Hillier L.,
 RA Nelson J., Spieth J., Ryan E., Andrews S., Geisel C., Layman D.,
 RA Du H., Ali J., Berghoff A., Jones K., Drone K., Cotton M., Joshi C.,
 RA Antoniou B., Zidanic M., Strong C., Sun H., Iamar H., Yordan C.,
 RA Ma P., Zhong J., Preston R., Vil D., Shekter M., Marero A., Shah A.,
 RA Swaby I.K., O'Shaughnessy A., Rodriguez M., Hoffman J., Tili S.,
 RA Granat S., Shohdy N., Hasegawa A., Hameed A., Lodhi M., Johnson A.,
 RA Chen E., Maria M., Martensen R., McCoy W.R.:
 RT "Sequence and analysis of chromosome 4 of the plant Arabidopsis
 RT thaliana";
 RL Nature 402:769-777(1999).
 RN [3]
 RP SEQUENCE OF 72-82; 96-110; 150-159; 178-211 AND 306-320.
 RA Schubert M., Peterson U., Funk C., Haus B., Schroeder W.P.,
 RA Kieselbach T.:
 RT "The chloroplast lumen from Arabidopsis thaliana";
 RL Submitted (JUL-2001) to the SWISS-PROT data bank.
 CC -1- SUBCELLULAR LOCATION: Chloroplast; within the thylakoid lumen.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S2C.
 CC -1- CAUTION: Ref.2 sequences differ from that shown due to erroneous
 CC gene model prediction. AT4G18370 and AT4G18375 were originally
 CC fused into a single gene.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its

CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; AF114386; AAF24060.1; -;
 CC EMBL; AL021710; CAA16717.1; ALT_SEQ.
 CC EMBL; AL161548; CAB78839.1; ALT_SEQ.
 CC MFROPS; S01.279; -;
 CC InterPro: IPR001940; ProteaseG2C.
 CC InterPro: IPR001254; Ser.protease_Try.
 CC Pfam: PF00089; trypsin; 1;
 CC PRINTS: PR00834; PROTEASES2C.
 CC Hydrolyase; Serine protease; Chloroplast; Thylakoid; Transit peptide.
 CC TRANSIT 1 26 CHLOROPLAST (POTENTIAL).
 CC TRANSIT 27 71 THYLAKOID.
 CC CHAIN 72 321 PROTEASE HHOA.
 CC DOMAIN 77 87 POLY-GLU.
 CC ACT_SITE 145 145 CHARGE RELAY SYSTEM (POTENTIAL).
 CC ACT_SITE 185 186 CHARGE RELAY SYSTEM (POTENTIAL).
 CC ACT_SITE 264 264 CHARGE RELAY SYSTEM (POTENTIAL).
 CC CONFLICT 40 40 R -> G (IN REF. 1).
 CC SEQUENCE 321 AA; 34691 MW; 66DB81E0BD27A7A7 CRC64;
 CC -----
 CC Query Match 8.8%; Score 84; DB 1; Length 321;
 CC Best Local Similarity 22.28; Pred. No. 0.88;
 CC Matches 48; Conservative 26; Mismatches 60; Indels 82; Gaps 11;
 CC -----
 CC 22 LTGRDKNQVEVOIVSTAAQIFLATCINGVCM-----TVYH----- 58
 CC 117 LTDEENKIKETG-----SGVWOKLGHIVNYHVIAKLATQFGLQRC 161
 CC 59 -----GAGTRTIASPKGVHVIOMYTNVDKWPAPQGSRLTCTCGSSDLYLVTRHAD 113
 CC 162 VSLVDKNGTR--FSKEGKIVCL--DPNDLAVLKIEIFGREINPVLGTSNDLRVGGSCF 217
 CC 114 VIPVRGRDSRG-----SLSPRISYUK-----GSSGGPLLCPA 148
 CC 218 AI-----GNPGYENTLTIGVSGIGREIPSPNGKISAEATQTDADINSNGSGGLDJSY 272
 CC 149 GHVNGIFRAAVCIIR--GVAKAVDF-IPVESLETIM 180
 CC 273 GHTIGV-NTATFTRRGSGVNAFPAIDTVVRIV 307
 CC -----
 CC RESULT 13
 CC AAMP_HUMAN STANDARD; PRT; 452 AA.
 CC ID Q13685;
 CC DT 15-JUL-1998 (Rel. 36, Created)
 CC DT 15-JUL-1998 (Rel. 36, Last sequence update)
 CC DT 28-FEB-2003 (Rel. 41, Last annotation update)
 CC DE Angio-associated migratory cell protein.
 CC GN AAMP.
 CC OS Homo sapiens (Human).
 CC CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC CC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 CC CC NCBI_TaxID=9606;
 CC RN [1]
 CC RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.
 CC RC 1:SSC=Brain;
 CC RX MEDLINE=95262124; PubMed=7743515;
 CC RA Beckner M.E., Krutzsch H.C., Stracke M.L., Williams S.T.,
 CC RA Gallardo J.A., Liotta L.A.;
 CC RT Identification of a new immunoglobulin superfamily protein expressed
 CC in blood vessels with a heparin-binding consensus sequence.*;
 CC RL Cancer Res. 55:2140-2149(1995).
 CC CC -1- FUNCTION: MAY HAVE A FUNCTION IN MIGRATING CELLS.
 CC CC -1- TISSUE SPECIFICITY: EXPRESSED IN BLOOD VESSELS. STRONGLY EXPRESSED
 CC IN ENDOTHELIAL CELLS, CYTOTROPHOBLASTS, AND POORLY DIFFERENTIATED
 CC COLON ADENOCARCINOMA CELLS FOUND IN LYMPHATICS.
 CC CC -1- SIMILARITY: Contains 8 WD repeats.
 CC -----

CC This SWISS-PROT entry is copyrighted. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announcement/>
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL: M95627; AAA69889.1; --
DR Genbank: U00018; AAMP.
DR MIM: 603488; --
DR GO: GO:0008201; Fibrinogen binding activity; TAS.
DR InterPro: IPR001680; WD40.
DR Pfam: PF00400; WD40; 8.
DR SMART: SM00320; WD40; 8.
DR PROSITE: PS00678; WD_REPEATS_1; 1.
DR PROSITE: PS50082; WD_REPEATS_2; 6.
DR PROSITE: PS50294; WD_REPEATS_REGION; 1.
KW Repeat; WD repeat.
FT DOMAIN 14 18 HEPARIN-BINDING (POTENTIAL).
FT DOMAIN 71 77 POLY-GLU.
FT REPEAT 107 138
FT REPEAT 150 180 WD 1.
FT REPEAT 190 220 WD 2.
FT REPEAT 231 261 WD 3.
FT REPEAT 275 306 WD 4.
FT REPEAT 333 363 WD 5.
FT REPEAT 374 404 WD 6.
FT REPEAT 416 446 WD 7.
FT REPEAT 452 485 WD 8.
SQ SEQUENCE 452 AA; DA1413D25E2B236C0 CRC64;

Query Match 8.6%; Score 82; DB 1; Length 452;

Best Local Similarity 25.3%; Pred. No. 2;

Matches 42; Conservative 13; Mismatches 47; Indels 64; Caps 9;

QY 54 WTIVHGAGRTIASPKGPVQKVTNVDKLVGPAQGSASL-----TPCTCGSSDLVY 108
DB 197 WMEWH-----PRAPVLACT-ADGNTWKKVNGNCKTFQGNCPAICGR----- 240
QY 109 TRHADVIVPRR-----GSRGSS-----LLSPKPSYLGKSGS--GPLCPA----- 148
DB 241 ----V-LPDKRAVGVYEDGTLRIWDLKQSG-IVLKGTHGHPGLTCVAANGDGLILT 295
QY 149 -----GHAVGIFR-----AAVCTRVAKAVDFPVSF, 176
DB 296 GSVDCQAKLVSAIGKVVGVFRPETVASQPSLSEGESESNSVSE, 341

RESULT 12

Y136_TREPA
ID Y136_TREPA STANDARD; PRT: 425 AA.
AC 081172;
DI 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DE 16-OCT-2001 (Rel. 40, Last annotation update)
DS Hypothetical lipoprotein TP0136 precursor.
GN TP0136.
OS Treponema pallidum.
OC Bacteria; Spirochaetes; Spirochaetales; Spirochaetaceae; Treponema.
OX NCBI_TaxID=160;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Nichols;
RX MEDLINE=98332770; PubMed=9655876;
RA Fraser C.M., Norris S.J., Weinstock G.M., White C., Sutton G.G.,
RA Dodson R., Guinn M., Hickey E.K., Clayton R., Ketchum K.A.,
RA Sodergren E., Hardham J.M., McLeod M.P., Salzberg S., Peterson J.,
RA Khalak H., Richardson D., Howell J.K., Chidambaram M., Utterback T.,
RA McDonald L., Artlich P., Bowman C., Cotton M.D., Fujii C., Garland S.,
RA Hatch B., Horst K., Roberts K., Sandusky M., Weidman J., Smith H.O.,
RA Venter J.C.;
*Complete genome sequence of Treponema pallidum, the syphilis

RT spirochete.";
RL Science 281:375-388(1998).
CC -!- SURCELLULAR LOCATION: Attached to the membrane by a lipid anchor
CC (Potential).
CC -!- SIMILARITY: BELONGS TO THE TP013X FAMILY OF LIPOPROTEINS.
CC -----
CC This SWISS-PROT entry is copyrighted. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announcement/>
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL: AE001199; AAC65137.1; ALT_INIT.
DR TIGR: TP0136; --
KW Hypothetical protein; Lipoprotein; Membrane; Signal;
KW Complete proteome.
FT SIGNAL 1 23 POTENTIAL.
FT CHAIN 24 485 HYPOTHETICAL LIPOPROTEIN TP0136.
FT LIPID 24 24 N-ACYL DIGLYCERIDE (POTENTIAL).
FT DOMAIN 164 178 GLY/SER-RICH.
FT DOMAIN 196 210 GLY/SER-RICH.
FT DOMAIN 253 267 GLY/SER-RICH.
FT DOMAIN 318 327 POLY-SER.
FT DOMAIN 444 447 POLY-SER.
SQ SEQUENCE 485 AA; 48984 MW; C7A4CEEDC7DC5CED CRC64;
Query Match 8.6%; Score 82; DB 1; Length 485;
Best Local Similarity 24.2%; Pred. No. 2,2;
Matches 44; Conservative 13; Mismatches 65; Indels 60; Gaps 8;
QY 23 TGRDKNQVEGEVQIVSTAAQTFLATCI--NGVWTVYHGAG---TRTIASPKGPVQKYT 77
DB 86 TSDSK-----KWSIAIDGNTFVLAQVPGVIGYKHCNVAGAGSSSTGTGTASPSTETCSQA 140
QY 78 NVYDKLVG-----WPAQGSRLTPCTC-----GSSDLVYTRHADVIP-----VR 118
DB 141 T-----LVGTSKPPVWLVPVGTGNNCGCGGGGGSSSSSSSSCHILVLPVGTGNNKCG 196
QY 119 RRGDSRGSLSPRLSYLK-----GSSGGPPLCPAGHA 151
DB 147 CGGGGGSSSSSSSSSSSSSSSIHKVENTDQFLDMGEGYVVT-KHLYTKNGSSSAGPAQCPCGGG 256
QY 152 VG 153
DB 257 GG 258
RESULT 13
DEG_ARATH
ID DEG_ARATH STANDARD; PRT: 437 AA.
AC 022609; O9LK85;
DI 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DE 28-FEB-2003 (Rel. 41, Last annotation update)
DS Protease Do-like 1, chloroplast precursor (EC 3.4.21.-).
GN DEGP1 OR DEGP OR A3G27525 OR K16N12.18.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A., AND CHARACTERIZATION.
RX MEDLINE=98175982; PubMed=9507020;
RA Itzhaki H., Naveh L., Lindahl M., Cook M., Adam Z.;
*Identification and characterization of DegP, a serine protease
RL associated with the luminal side of the thylakoid membrane.";
RL J. Biol. Chem. 273:7094-7098(1998).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Columbia;

Dd		210 PK--NKLRIPIVGVNSADLLVGQKVFALGNPFGLDHLTLTGIVISGLRRRHS--SAATGRPI 266
QY		134 SYL-----KGSSGGPFLLCAGHAGVIFRAAVCTRGAKAYDF-IPVESL 176 :
Dd		266 QDVIOQTAAINPGSGGPLEDDSSGLIGINTAIYSFGASSGVGFSPVDTV 317 :
RESULT 14		
ID	CAC3_BOVIN	STANDARD:
AC	P58005:	FRT: 253 AA.
DT	01-NOV-1988 (Rel. 09, Created)	
DI	15-DEC-1998 (Rel. 37, Last sequence update)	
DE	15-SEP-2003 (Rel. 42, Last annotation update)	
DE	Proteinase E precursor [Procarboxypeptidase A complex component	
CE	(II) (Procarboxypeptidase A-S6 subunit III) (PROCPA-S6 III).	
CS	Bos taurus (Bovine)	
CC	Eukaryota; Metazoa; Chordata; Cranialia; Vertebrata; Euteleostomi;	
OC	Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;	
OC	Bovidae; Bovinae; Bos.	
CX	NCBM_LaxID=9913;	
PN	[1]	
KP	SOURCE OF 1-25.	
RX	MEDLINE=F91099520; PubMed=2269366;	
RX	Pascual K., Vendrell J.J., Aviles F.X., Bonicel J., Wicker C.,	
KA	Puigserver A.;	
KT	"Autolysis of proproteinase E in bovine procarboxypeptidase A ternary	
RL	complex gives rise to subunit I11.";	
PL	FEBS Lett. 277:37-4;(1990).	
RN	[2].	
RN	SEQUENCE OF 14-253, AND DISULFIDE BONDS.	
RX	MEDLINE=86220198; Pubmed=3519215;	
RX	Venot N., Sciaky M., Puigserver A., Desnuelle P., Laurent G.;	
RC	Amino acid sequence and disulfide bridges of subunit I11, a	
RJ	defective endopeptidase present in the bovine pancreatic 6 S	
RJ	procarboxypeptidase A complex.";	
RI	Eur. J. Biochem. 157:91-99(1986).	
RN	[3].	
RP	X-RAY CRYSTALLOGRAPHY (1.7 ANGSTROMS).	
RX	MEDLINE=94222022; PubMed=8168476;	
RA	Pignol D., Gaboriau C., Michon L., Kerfelec B., Chapus C.,	
RA	Fontecilla-Camps J.C.;	
RI	"Crystal structure of bovine procarboxypeptidase A-S6 subunit I11, a	
RJ	highly structured truncated zymogen E.";	
RC	EMBO J. 13:1763-1771(1994).	
CC	-I- FUNCTION: DEFECTIVE ELASTASE-LIKE SERINE PROTEASE. DOES NOT SEEM	
CC	TO HAVE A PROTEASE ACTIVITY. ITS LIKELY FUNCTION IS TO PROTECT	
CC	PROCARBOXYPEPTIDASE A AGAINST DENATURATION IN THE ACIDIC	
CC	ENVIRONMENT OF THE RUMINANT DUODENUM.	
CC	-I- SUBUNIT: HETEROTRIMER OF SUBUNIT I11; CARBOXYPEPTIDASE A AND	
CC	CHYMOTRYPSINOGEN C.	
CC	-I- SUBCELLULAR LOCATION: Extracellular.	
CC	-I- TISSUE SPECIFICITY: Pancreas.	
CC	-I- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.	
DR	PDB: IFON; 14-OCT-96.	
DR	PDB: IPYT; 27-JAN-97.	
DR	MEROPS: S01.983.	
DR	InterPro: IPR001314; Chymotrypsin.	
DR	InterPro: IPR001254; Ser_protease_Iry.	
DR	Prints: PF00089; trypsin; 1.	
DR	SMART: SM00020; Tryp_SPC; 1.	*
DR	PROSITE: PS00240; TRYPSIN_DOM; 1.	
DR	PROSITE: PS00134; TRYPSIN_HIS; 1.	
DR	PROSITE: PS00135; TRYPSIN_SER; 1.	
KW	Serine protease homolog; Pancreas; Digestion; 3D-structure.	
FT	PROPEP 1 11	ACTIVATION PEPTIDE.
FT	CHAIN 12 253	PROPEPTINASE E.
FT	DISULFID 41 57	
FT	DISULFID 100 103	
FT	DISULFID 140 206	
FT	DISULFID 171 187	
FT	DISULFID 196 227	

Result No.	Score	Query Match	Length	DR	ID	Description
1	94.3	99.0	3011	1	GNWVC3	genome polyprotein
2	93.9	98.5	3011	1	S40770	genome polyprotein
3	92.7	97.3	3011	1	GNWVCH	genome polyprotein
4	89.2	93.6	3010	1	GNWVTW	genome polyprotein
5	89.1	93.5	3010	1	GNWVTC	genome polyprotein
6	88.7	93.1	3010	1	GNWVCJ	genome polyprotein
7	88.4	92.8	3010	1	A45573	genome polyprotein
8	86.4	90.7	3010	1	S18030	genome polyprotein
9	80.4	84.4	3014	1	JC5620	genome polyprotein
10	71.4	74.9	3033	1	GNWVJ8	genome polyprotein
11	71.2	74.7	3033	1	JQ1303	genome polyprotein
12	267.5	28.1	3025	1	T08841	polyprotein - dsur
13	255.5	26.6	2970	2	T08839	polyprotein - marm
14	87	9.1	209	2	H8144	probable aromatic
15	82	8.6	452	2	I39383	angio-associated m
16	82	8.6	495	2	B71360	hypothetical prote
17	91.5	8.6	476	2	T48199	heat shock transer
18	78.5	8.2	620	2	F83976	cytochrome c oxida
19	78.5	8.2	981	2	T18234	beta transducin ho
20	78	8.2	239	2	H89566	serine proteinase
21	77.5	8.1	398	2	B71284	probable periplasm
22	77	8.1	904	2	AB4212	hypothetical prote
23	76.5	8.0	270	2	T06118	hypothetical prote
24	76.5	8.0	868	2	H81775	aconitate hydratase
25	75	7.9	240	1	CP60A3	procarboxypeptidas
26	74.5	7.8	415	2	S79401	zona pellucida gly
27	74.5	7.8	735	2	S23441	hypothetical prote
28	74.5	7.8	868	2	C81820	aconitate hydratase
29	74.5	7.8	911	2	JN0821	transferrin-binding


```
1086 GSKTAGKGPITQMYTNVQDLVGVHPAPGASMPCTCGSSDLVLTIRHADVVPVRR 1145
QY 121 GDSRGSLLSPRPISYLKSSGGPLLCAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
DB 1146 GDSRGSLLSPRPISYLKSSGGPLLCAGHAGVIFRAAVCTRGVAKAVDFIPVESMETTM 1205
QY 181 RS 182
DB 1206 RS 1207

RESULT 7
genome polyprotein - hepatitis C virus (strain J1)
N:Contains: capsid protein C; envelope protein M; hepatitis virus (EC 3.4.21.98) (nonstru
protein NS4a; nonstructural protein NS4b; nonstructural protein NS5
C:Species: hepatitis C virus
C:Date: 19-May-2000 #sequence_revision 19-May-2000 #text_change 19-Jan-2001
C:Accession: A45573
R:Tanaka, T.; Kato, N.; Nakagawa, M.; Ootsuyama, Y.; Cho, M.J.; Nakazawa, F.; Hijikata,
Virus Res. 23, 39-53, 1992
A:Title: Molecular cloning of hepatitis C virus genome from a single Japanese carrier: s
A:Reference number: A45573; MUID:92295714; PMID:1335627
A:Accession: A45573
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-3010 <TAN>
A:Cross-references: GB:D01171; NID:q221612; PIDN:BAA01943.1; PID:q221613
A:Experimental source: HCV-JT
A:Note: sequence extracted from NCBI backbone (NCBIN:106206, NCBIPI:106207)
C:Superfamily: hepatitis C virus genome polyprotein
C:Keywords: ATP; glycoprotein; hydrolase; nucleotide binding; P-loop; polyprotein; scri
F:2-115/Product: capsid protein C #status predicted <CPC>
F:116-191/Product: envelope protein M #status predicted <EPM>
F:192-389/Product: major envelope protein E #status predicted <MEP>
F:390-729/Product: nonstructural protein NS1 #status predicted <NS1>
F:730-1006/Product: nonstructural protein NS2 #status predicted <NS2>
F:1007-1615/Product: hepatitis virus #status predicted <NS3>
F:1230-1237/Region: nucleotide-binding motif A (P-loop)
F:1316-1319/Region: DEXH motif
F:1616-1862/Product: nonstructural protein NS4a #status predicted <N4A>
F:1863-2013/Product: nonstructural protein NS4b #status predicted <N4B>
F:2014-3010/Product: nonstructural protein NS5 #status predicted <NS5>

Query Match 92.8% Score 884 DB 1 Length 3010
Best Local Similarity 88.5% Pred. No. 2.3e-75
Matches 161 Conservative 15 Mismatches 6 Indels 0 Gaps 0
QY 1 MAPITAAQOTRGLGCIITSLTGDRKNQVGEVQIVSTAAQIFATCINXGVCTVYHGA 60
DB 1026 LAPITAAQOTRGLGCIITSLTGDRKNQVGEVQIVSTAAQIFATCINXGVCTVYHGA 1085
QY 61 GTRITASPKGPVQMYTNVQDLVGVHPAPGASMPCTCGSSDLVLTIRHADVVPVRR 120
DB 1086 GSKTAGKGPITQMYTNVQDLVGVHPAPGASMPCTCGSSDLVLTIRHADVVPVRR 1145
QY 121 GDSRGSLLSPRPISYLKSSGGPLLCAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
DB 1146 GDSRGSLLSPRPISYLKSSGGPLLCAGHAGVIFRAAVCTRGVAKAVDFIPVESMETTM 1205
QY 181 RS 182
DB 1206 RS 1207

RESULT 8
S18030
genome polyprotein - hepatitis C virus (isolate J1)
N:Contains: capsid protein C; envelope protein M; hepatitis virus (EC 3.4.21.98) (nonstru
protein NS4a; nonstructural protein NS4b; nonstructural protein NS5
C:Species: hepatitis C virus
C:Date: 19-May-2000 #sequence_revision 19-May-2000 #text_change 19-Jan-2001
C:Accession: J18030
R:Chamberlain, R.W.; Adams, N.J.; Taylor, L.A.; Simmonds, P.; Elliott, R.M.
Biochem. Biophys. Res. Commun. 236, 44-49, 1997
A:Title: The complete coding sequence of hepatitis C virus genotype 5a, the predomina
A:Reference number: J18030; MUID:97366593; PMID:9223423
```

```
A:Variety: isolate J1
C:Date: 19-May-2000 #sequence_revision 19-May-2000 #text_change 23-Mar-2001
C:Accession: S18030; S33570; A48332; S18029
R:Honda, M.; Kaneko, S.; Masashi, U.; Kobayashi, K.; Murakami, S.
submitted to the EMBL Data Library, September 1991
A:Description: A whole genome of hepatitis C virus cDNA was isolated from a single i
A:Reference number: S18028
A:Accession: S18030
A:Molecule type: genomic RNA
A:Residues: 1-3010 <HON>
A:Cross-references: EMBL:X61596; NID:q59478; PIDN:CAA4793.1; PID:q59479
A:Experimental source: isolate J1 from an individual
R:Honda, M.; Kaneko, S.; Uemura, M.; Kobayashi, K.; Murakami, S.
Arch. Virol. 128, 163-169, 1993
A:Title: Sequence analysis of putative structural regions of hepatitis C virus isolat
A:Reference number: A48332; MUID:9119270; PMID:8380322
A:Accession: S33570
A:Molecule type: genomic RNA
A:Residues: 1-547, 'T', '549-621', 'V', '623-624', 'S', '626-652', 'DL', '655-761', 'T', '763-782' <HOW>
A:Cross-references: EMBL:X61591
A:Note: this sequence is inconsistent with the nucleotide translation
as Trp, and TTC for residue 771 as Ser
A:Note: sequence extracted from NCBI backbone (NCBIN:121747, NCBIPI:121748)
C:Superfamily: hepatitis C virus genome polyprotein
C:Keywords: ATP; glycoprotein; hydrolase; nucleotide binding; P-loop; polyprotein; s
F:2-115/Product: capsid protein C #status predicted <CPC>
F:116-191/Product: envelope protein M #status predicted <EPM>
F:192-389/Product: major envelope protein E #status predicted <MEP>
F:390-729/Product: nonstructural protein NS1 #status predicted <NS1>
F:730-1006/Product: nonstructural protein NS2 #status predicted <NS2>
F:1007-1615/Product: hepatitis virus #status predicted <NS3>
F:1230-1237/Region: nucleotide-binding motif A (P-loop)
F:1316-1319/Region: DEXH motif
F:1616-1862/Product: nonstructural protein NS4a #status predicted <N4A>
F:1863-2013/Product: nonstructural protein NS4b #status predicted <N4B>
F:2014-3010/Product: nonstructural protein NS5 #status predicted <NS5>
F:196,209,234,250,305,417,423,448,532,540,556,576,623,643,645/Binding site: carbohydrate
Query Match 90.7% Score 864 DB 1 Length 3010
Best Local Similarity 86.8% Pred. No. 1.9e-73
Matches 158 Conservative 15 Mismatches 9 Indels 0 Gaps 0
QY 1 MAPITAAQOTRGLGCIITSLTGDRKNQVGEVQIVSTAAQIFATCINXGVCTVYHGA 60
DB 1026 LAPITAAQOTRGLGCIITSLTGDRKNQVGEVQIVSTAAQIFATCINXGVCTVYHGA 1085
QY 61 GTRITASPKGPVQMYTNVQDLVGVHPAPGASMPCTCGSSDLVLTIRHADVVPVRR 120
DB 1086 GSKTAGKGPITQMYTNVQDLVGVHPAPGASMPCTCGSSDLVLTIRHADVVPVRR 1145
QY 121 GDSRGSLLSPRPISYLKSSGGPLLCAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
DB 1146 GDSRGSLLSPRPISYLKSSGGPLLCAGHAGVIFRAAVCTRGVAKAVDFIPVESMETTM 1205
QY 181 RS 182
DB 1206 RS 1207

RESULT 9
JC5620
genome polyprotein - hepatitis C virus (isolate EUH1480)
N:Contains: capsid protein C; envelope protein M; hepatitis virus (EC 3.4.21.98) (nonstru
protein NS4a; nonstructural protein NS4b; nonstructural protein NS5
C:Species: hepatitis C virus
C:Date: 19-May-2000 #sequence_revision 19-May-2000 #text_change 19-Jan-2001
C:Accession: JC5620
R:Chamberlain, R.W.; Adams, N.J.; Taylor, L.A.; Simmonds, P.; Elliott, R.M.
Biochem. Biophys. Res. Commun. 236, 44-49, 1997
A:Title: The complete coding sequence of hepatitis C virus genotype 5a, the predomina
A:Reference number: JC5620; MUID:97366593; PMID:9223423
```

A:Accession: J05620
A:Molecule type: mRNA
A:Residues: 1-3014 <CH>
A:Cross-references: GB:D10388; GB:D01221; NID:g221608; PIDN:BA01761.1; PID:g221609
A:Experimental source: isolate F-012
A:Note: The translation of the nucleotide sequence is not complete in this paper
C:Superfamily: hepatitis C virus genome polyprotein
C:Keywords: ATP; glycoprotein; hydrolyase; nucleotide binding; P-loop; polyprotein; serine
F:116-191/Product: capsid protein C #status predicted <CPC>
F:192-389/Product: major envelope protein E #status predicted <EPM>
F:384-408/Product: hypervariable #status predicted
F:390-730/Product: nonstructural protein NS1 #status predicted <NS1>
F:731-1007/Product: nonstructural protein NS2 #status predicted <NS2>
F:1008-1616/Product: hepatitis C virus genome polyprotein NS3 #status predicted <NS3>
F:1231-1238/Region: nucleotide-binding motif A (P-loop)
F:1313-1318/Region: DEXH motif
F:1317-1320/Region: DEXH motif
F:1617-1863/Product: nonstructural protein NS4a #status predicted <NS4a>
F:1864-2014/Product: nonstructural protein NS4b #status predicted <NS4b>
F:2015-3014/Product: nonstructural protein NS5 #status predicted <NS5>
F:2210-2249/Region: interferon sensitivity determining #status predicted

Query Match 84.4% Score 804; DB 1; Length 3014;
Best Local Similarity 79.1%; Pred. No. 9.5e-68;
Matches 144; Conservative 22; Mismatches 16; Indels 0; Gaps 0;

QY 1 MAPITAYAAQQTGGLGCIITSLTGDRKNQVEGEVIVSTAAQTFLATCINGVCWTVYHGA 60
DB 1027 LAPITAYAAQQTGGLGCIITSLTGDRKNQVEGEVIVSTAAQTFLATCINGVCWTVYHGA 1084

QY 61 GRTIASPKGPIVOMYTNVDKLVGNPAPQGSRSITPCTCGSSDLVYVTRHADVPIVRR 120
DB 1087 GSKLAGKGPVOMYTNVDKLVGNPAPQGSRSITPCTCGSSDLVYVTRHADVPIVRR 1146

QY 121 GDSRSLSPRISYLGSSGGLPCAGHAGVIFRAAVCTRGVAKAVDFIPVESLETIM 180
DB 1147 GDSRSLSPRISYLGSSGGLPCAGHAGVIFRAAVCTRGVAKAVDFIPVESLETIM 1206

QY 161 RS 162
DB 1207 RS 1208

RESULT 10
GNWJ8
genome polyprotein - hepatitis C virus (strain HC-J8)
N:Contains: capsid protein C; envelope protein M; hepatitis C virus (strain HC-J8) (nonstructural protein NS4a; nonstructural protein NS4b; nonstructural protein NS5)
C:Species: Hepatitis C virus
C:Date: 31-Dec-1992 #sequence revision 31-Dec-1992 #text_change 19-Jan-2001
C:Accession: A40250; PC0357; PC0559
R:Okamoto, H.; Kunita, K.; Okada, S.; Yamamoto, K.; Iizuka, H.; Tanaka, T.; Fukuda, S.; Virology 188, 331-341, 1992
A:Title: Full-length sequence of a hepatitis C virus genome having poor homology to reported reference number: A40250; MUID:92230232; PMID:1314459
A:Accession: A40250
A:Molecule type: genomic RNA
A:Residues: 1-3033 <OK>
A:Cross-references: GB:D01221; NID:g221608; PIDN:BA01761.1; PID:g221609
R:Chan, S.W.; McMahon, F.J.; Holmes, E.C.; Dow, B.; Penhale, J.F.; Follett, E.; Yap, P.H. J. Gen. Virol. 73, 1131-1141, 1992
A:Title: Analysis of a new hepatitis C virus type and its phylogenetic relationship to other hepatitis C virus types
A:Accession: PC0397
A:Molecule type: genomic RNA
A:Residues: 2578-2754 <CH>
A:Cross-references: DDB:210134
A:Experimental source: isolate F-012
R:Kato, N.; Ootsuyama, Y.; Okoshi, S.; Nakazawa, T.; Mori, S.; Hijikata, M.; Shimotohno, K. Biochem. Biophys. Res. Commun. 181, 279-285, 1991
A:Title: Distribution of pluri HCV types in Japan.
A:Reference number: PC0554; MUID:92068204; PMID:1720409

A:Accession: PQ0559
A:Molecule type: mRNA
A:Residues: 2678-2729 <KAT>
A:Cross-references: GB:D10562; GB:D090518; NID:g221523; PIDN:BA01418.1; PID:g221524
C:Superfamily: hepatitis C virus genome polyprotein
C:Keywords: ATP; capsid protein; envelope protein; glycoprotein; hydrolyase; nonstructural protein NS4a; nonstructural protein NS4b; nonstructural protein NS5
F:115-191/Product: capsid protein C #status predicted <CPC>
F:116-191/Product: major envelope protein E #status predicted <EPM>
F:192-389/Product: major envelope protein E #status predicted <EPM>
F:390-730/Product: nonstructural protein NS1 #status predicted <NS1>
F:734-1010/Product: nonstructural protein NS2 #status predicted <NS2>
F:1011-1619/Product: hepatitis C virus genome polyprotein NS3 #status predicted <NS3>
F:1234-1244/Region: nucleotide-binding motif A (P-loop)
F:1316-1322/Region: nucleotide-binding motif B
F:1320-1323/Region: DEXH motif
F:1620-1866/Product: nonstructural protein NS4a #status predicted <NS4a>
F:1867-2017/Product: nonstructural protein NS4b #status predicted <NS4b>
F:2018-3033/Product: nonstructural protein NS5 #status predicted <NS5>
F:2195-205,233,299,305,417,423,430,448,477,534,542,558,578,627,649,1091,1217,1259,21

Query Match 74.9% Score 714; DB 1; Length 3033;
Best Local Similarity 70.9%; Pred. No. 3.5e-59;
Matches 129; Conservative 26; Mismatches 27; Indels 0; Gaps 0;

QY 1 MAPITAYAAQQTGGLGCIITSLTGDRKNQVEGEVIVSTAAQTFLATCINGVCWTVYHGA 60
DB 1030 LAPITAYAAQQTGGLGCIITSLTGDRKNQVEGEVIVSTAAQTFLATCINGVCWTVYHGA 1089

QY 61 GRTIASPKGPIVOMYTNVDKLVGNPAPQGSRSITPCTCGSSDLVYVTRHADVPIVRR 120
DB 1090 GRTIASPKGPIVOMYTNVDKLVGNPAPQGSRSITPCTCGSSDLVYVTRHADVPIVRR 1149

QY 121 GDSRSLSPRISYLGSSGGLPCAGHAGVIFRAAVCTRGVAKAVDFIPVESLETIM 180
DB 1150 GDSRSLSPRISYLGSSGGLPCAGHAGVIFRAAVCTRGVAKAVDFIPVESLETIM 1209

QY 181 RS 182
DB 1210 RS 1211

RESULT 11
JQ1303
genome polyprotein - hepatitis C virus (isolate HC-J6)
N:Contains: capsid protein C; envelope protein M; hepatitis C virus (isolate HC-J6) (nonstructural protein NS4a; nonstructural protein NS4b; nonstructural protein NS5)
C:Species: hepatitis C virus
C:Date: 19-May-2000 #sequence revision 19-May-2000 #text_change 17-Nov-2000
C:Accession: JQ1303
R:Okamoto, H.; Okada, S.; Sugiyama, Y.; Kunita, K.; Iizuka, H.; Machida, A.; Miyake, J. Gen. Virol. 72, 2697-2704, 1991
A:Title: Nucleotide sequence of the genomic RNA of hepatitis C virus isolated from a patient with chronic hepatitis C
A:Reference number: JQ1303; MUID:92044440; PMID:1658196
A:Accession: JQ1303
A:Molecule type: genomic RNA
A:Residues: 1-3033 <OK>
A:Cross-references: GB:D090518; NID:g221650; PIDN:BA00792.1; PID:g221651
A:Experimental source: isolate HC-J6 from a Japanese individual
C:Superfamily: hepatitis C virus genome polyprotein
C:Keywords: ATP; glycoprotein; hydrolyase; P-loop; polyprotein; serine proteinase;
F:116-191/Product: capsid protein C #status predicted <CPC>
F:192-389/Product: major envelope protein E #status predicted <EPM>
F:390-733/Product: nonstructural protein NS1 #status predicted <NS1>
F:734-1010/Product: nonstructural protein NS2 #status predicted <NS2>
F:1011-1619/Product: hepatitis C virus genome polyprotein NS3 #status predicted <NS3>
F:1316-1321/Region: nucleotide-binding motif B
F:1320-1323/Region: DEXH motif
F:1620-1866/Product: nonstructural protein NS4a #status predicted <NS4a>
F:1867-2017/Product: nonstructural protein NS4b #status predicted <NS4b>
F:2018-3033/Product: nonstructural protein NS5 #status predicted <NS5>
F:196,209,234,305,325,417,423,430,448,477,534,542,558,578,627,649,1091,1217,1259,21

Query Match 74.7% Score 712; DB 2; Length 1033;
Best Local Similarity 64.8%; Pred. No. 54e-53;
Matches 127; Conservative 29; Mismatches 26; Indels 0; Gaps 0;
QY 1 KAPITAYAOQTGRLGCIITSLTGRKNOVEGEVQIVSTAAQTFLATCINGVCWVYHGA 60
DB 2030 KAPITAYAOQTGRLGCIITSLTGRKNOVEGEVQIVSTAAQTFLATCINGVCWVYHGA 1099
QY 61 CTRTIASPKGPVIONYTNVDKDLVGPAPQGSRLTFTCGSSDLVLTTRHADVPVRRR 120
DB 1090 GNTIAGSRPVTOMYSSAGDVLGVPSPGTSKSEPCCGAVDLYLTNRADVPARRR 1149
QY 121 GDSRGLSPRPISYKSGSGGGLPCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLET 180
DB 1150 GDRKGLSPRPISYKSGSGGGLPCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLET 1209
QY 181 RS 182
DB 1210 RS 1211
RESULT 12
T08841
Polyprotein - douroucouli hepatitis GB virus A
C:Species: douroucouli hepatitis GB virus A
C:Date: 20-Sep-1999 #sequence_revision 20-Sep-1999 #text_change 17-Nov-2000
C:Accession: T08841
R:Erker, J.C.; Desai, S.M.; Leary, T.P.; Chalmers, M.L.; Montes, C.C.; Mushahwar, I.K.
J. Gen. Virol. 79, 41-45, 1998
A:Title: Genomic analysis of two GB virus A variants isolated from captive monkeys.
A:Reference number: Z16486; MUID:98120818; PMID:9460920
A:Accession: T08841
A:Molecule type: mRNA
A:Status: translated from GB/EMBL/DBJ
A:Residues: 1-3005 <ERK>
A:Cross-references: EMBL:AF023425; NID:g2828599; PIDN:AAC40502.1; PID:g2828630
A:Superfamily: hepatitis C virus genome polyprotein
C:Keywords: polyprotein
Query Match 28.1% Score 267.5; DB 2; Length 3005;
Best Local Similarity 33.1%; Pred. No. 1e-16;
Matches 60; Conservative 32; Mismatches 75; Indels 11; Gaps 4;
QY 2 APTTAYAOQTGRLGCIITSLTGRKNOVEGEVQIVSTAAQTFLATCINGVCWVYHGA 61
DB 579 APVVV-MORGLGPFSSVVKTSMEGRDEREHEGSIIVLTSTIRSMGICVNGVMYITFHGSN 1037
QY 52 IRTIASPKGPVIONYTNVDKDLVGPAPQGSRLTFTCGSSDLVLTTRHADVPVRRR 121
DB 1038 ARTIAGPVPVNCRWSPSDVAVYPTSGASCTHEPKCTQSVWCTRN--DGAICGRL 1095
QY 122 DSRGSLSPRPISYKSGSGGGLPCPAGHAGVIFRAAVCTRGVAKAVDFIPVE 174
DB 1096 SKVLVDLPTTISDFRSGSGSPICDEGHVGMZ-VSVLHROGVKTVGVYKFWMLFKS 1154
QY 175 S 175
DB 1155 S 1155
RESULT 13
T08839
polyprotein - marmoset hepatitis GB virus A
C:Species: marmoset hepatitis GB virus A
C:Date: 23-Jul-1999 #sequence_revision 23-Jul-1999 #text_change 17-Nov-2000
C:Accession: T08839
R:Erker, J.C.; Desai, S.M.; Leary, T.P.; Chalmers, M.L.; Montes, C.C.; Mushahwar, I.K.
J. Gen. Virol. 79, 41-45, 1998
A:Title: Genomic analysis of two GB virus A variants isolated from captive monkeys.
A:Reference number: Z16486; MUID:98120818; PMID:9460920
A:Accession: T08839
A:Status: translated from GB/EMBL/DBJ
A:Molecule type: genomic RNA

A:Residues: 1-2970 <ERK>
A:Cross-references: EMBL:AF023424; NID:g2828597; PIDN:AAC40501.1; PID:g2828598
C:Superfamily: hepatitis C virus genome polyprotein
C:Keywords: polyprotein
Query Match 26.8% Score 255.5; DB 2; Length 2970;
Best Local Similarity 30.1%; Pred. No. 1.4e-15;
Matches 59; Conservative 36; Mismatches 66; Indels 33; Gaps 6;
QY 2 APTTAYAOQTGRLGCIITSLTGRKNOVEGEVQIVSTAAQTFLATCINGVCWVYHGA 61
DB 970 APVVVH-HHGKGFVGVKISMTGDETEHGVNVVVLGISTIRSMGTCVNGVMYITFHGSN 1028
QY 62 TRTIASPKGPVIONYTNVDKDLVGPAPQGSRLTFTCGSSDLVLTTRHADVPVRRR 121
DB 1029 ARTIAGPVPVNCRWSPSDVAVYPLVPGAKCLEPKCQCPQGVWVI-----RN 1077
QY 122 DSRGSLSPRPISYKSGSGGGLPCPAGHAGVIFRAAVCTRG----- 163
DB 1078 D--GALCHCTGRTVELDPAELCDFRSGSPILCDGHAAGML-ISVLRGSRVGTGR 1134
QY 164 VAKAVDFIPVESLETT 179
DB 1135 YTKPWETLPREATHY 1150
RESULT 14
H83144
Probable aromatic acid decarboxylase PA4019 [imported] - Pseudomonas aeruginosa (str)
C:Species: Pseudomonas aeruginosa
C:Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 31-Dec-2000
C:Accession: H83144
R:Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warren, P.; Hickey, M.J.
adman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.; Latbig, K.;
Lory, S.; Olson, M.V.
Nature 406, 959-964, 2000
A:Title: Complete genome sequence of Pseudomonas aeruginosa PA01, an opportunistic
A:Reference number: A82950; MUID:20437337; PMID:10964043
A:Accession: H83144
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-209 <STO>
A:Cross-references: GB:AE004816; GB:AE004091; NID:g9950200; PIDN:AAG07406.1; GSPDB:
A:Experimental source: strain PA01
C:Genetics:
A:Gene: PA4019
C:Superfamily: dedF protein
Query Match 9.1% Score 87; DB 2; Length 209;
Best Local Similarity 26.2%; Pred. No. 0.69;
Matches 55; Conservative 18; Mismatches 61; Indels 76; Gaps 12;
QY 8 AQQTGRLGCIITSLTGRKNOVEGEVQIVSTAAQTFLATCINGVCWVYHGA 66
DB 17 AQYGLRLDCLV-----QGEREVHFLISKRAQLVMAI-----ETDVA 53
QY 67 SPKGP-----VIQMYTNVDKDLVGPAPQGSRLTFTCGSSDL 105
DB 54 LPAPQMAQAFLEYCGAAGQIRVFGND-----WMAAPPASGSSAPNMMVICPSTGIL 108
QY 106 -----YLVTRHADVPVRRRGRSGSLSPR--PIS-----YKSGSGGGLPCPA 148
DB 109 SAVATGACNNLIFRAADVALKER-----RPLVLPREAPFSSIHLENMMLKLSNLAJAVILPA 164
QY 149 GHAVGIFRAAVCTRGVAKAVDFIPVESLET 178
DB 165 --APGFYHQ---POSVEDLYVFEVVARILNT 189
RESULT 15
I39383
angio-associated migratory cell protein - human
C:Species: Homo sapiens (man)

C:Date: 06-Sep-1996 #sequence_revision 06-Sep-1996 #text_change 21-Jul-2000
 C:Accession: 139383
 R:Beckner, N.E.; Krutzsch, H.C.; Stracke, N.L.; Williams, S.T.; Gallardo, J.A.; Liotta, L.
 Cancer Res. 55, 2140-2145, 1995
 A:Title: Identification of a new immunoglobulin superfamily protein expressed in blood v
 A:Reference number: 139383; PMID:95262124; PMID:773515
 A:Accession: 139383
 A>Status: preliminary; translated from GR/EMBL/DDBJ
 A:Molecule type: mRNA
 A:Residues: 1-452 <RES>
 A:Cross-references: GR:M95627; NID:q870802; PID:AAA68889.1; PID:q870803
 C:Genetics:
 A:Gene: GDR:AAMP
 A:Cross-references: GDB:457393
 A:Map position: 14q32.1-14q32.1
 C:Superfamily: unassigned WD repeat proteins; WD repeat homology
 F:148-181/Domain: WD repeat homology <WD1>
 F:414-447/Domain: WD repeat homology <WD2>

Query Match 8.64; Score 82; DB 2; Length 452;
 Best Local Similarity 25.3%; Pred. No. 57
 Matches 42; Conservative 13; Mismatches 47; Indels 44; Gaps 9;
 QY 54 WTVEGASTRTIASPKGPVIGWYTNVDRKLYGVWHPAQGSRSL-----TPTCGSSDLYLV 108
 DB 197 KWEWH-----PRAPVLLAST-ADGNTWKKVYPNGDKX-FOGPNCFATCGK----- 240
 QY 109 TRHADVIPRRR---GDSRGS-----LSRPPISTYKQSSG--GPIILGFA----- 148
 DB 241 -----VLEPGKRAVVGVEGTIRINDELKGGSPHVKVKTGTHGPGPLTCVAANQDSILL 295
 QY 149 -----CHAVGIER-----AAVCTRGVAKAVDFIPVESL 176
 DB 296 GSVDGQAKLVNATTKGVVPRPPTVVASQPSLGGEGESFSNSVESL 341

Search completed: September 27, 2003, 12:22:12
 Job time : 44 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - nucleic search, using frame_plus_p2n model

Run on: September 27, 2003, 12:28:45 : Search time 2332 Seconds
(without alignments)
1981.817 Million c.c.: updates/sec

Title: US-09-965-594-1
Perfect score: 953
Sequence: 1 MAPITAYAAQFRGLLSC-IT.....GVAXAVDFIPVESLETIMRS 182

Scoring table: BLOSUM62
Xgapop 10.0, Xgapext 0.5
Ygapop 10.0, Ygapext 0.5
Fgapop 6.0, Fgapext 7.0
Delop 6.0, Delext 7.0

Searched: 22781192 seqs, 12152235056 residues

Total number of hits satisfying chosen parameters: 45362744

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 04
Maximum Match 994
Listing first 45 summaries

Command line parameters: -DEV-xih
-O=/cgn2.1/USPTO.spool/US09965594/runat_26092003.1641433.18769/aff_query_fastal.1.327
-DB=EST -QFMT=fastop -SUFFIX=first -MINMATCH=9.1 -LOCAL=0 -LOCAL=0
-UNITS=bits -START=1 -END=1 -MATRIX=blosum62 -TRANS=human40.cdi -LIST=45
-DOCALIGN=200 -THR_SCORE=pct -THR_MAX=99.9 -THR_MIN=0 -ALIGN=15 -MCDE=LOCAL
-OUTFMT=pco -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000
-USER=US09965594 -CGN 1.1.2810 -runat_26092003.1641433.18769 -NCPU=6 -ICPU=3
-NO_MMAP -LARGEQUERY -NEG_SCORES=0 -WAIT_DSPBLOCK=100 -LONGLOG
-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -Fgapop=6
-Fgapext=7 -YGAPOP=10 -YGAPEXT=9.5 -DELOP=5 -DELEXT=7

Database :

EST: *
1: cm_estba: *
2: cm_estcum: *
3: cm_estin: *
4: cm_estnu: *
5: cm_estov: *
6: cm_estpl: *
7: cm_estro: *
8: cm_hic: *
9: gb_est1: *
10: gb_est2: *
11: gb_hic: *
12: gb_est3: *
13: gb_est4: *
14: gb_est5: *
15: cm_estfun: *
16: cm_estom: *
17: em_gss_hum: *
18: em_gss_inv: *
19: em_gss_pln: *
20: em_gss_vrt: *
21: em_gss_fun: *
22: em_gss_mam: *
23: em_gss_mus: *
24: em_gss_pro: *
25: em_gss_rod: *
26: em_gss_phg: *
27: em_gss_vrl: *
28: gb_gssl: *

29: gb_gss2: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
C 1	106	11.1	984	10	BF304699	BF304699 601888252
C 2	103.5	10.9	1199	13	B0892487	B0892487 AGENCOURT
C 3	101	10.6	515	14	CA023748	CA023748 H247817
C 4	101	10.6	583	12	BM374064	BM374064 Ema03_SO
C 5	99	10.4	515	12	BJ001625	BJ001625 BJO01625
C 6	99	10.4	643	12	BJ024121	BJ024121 BJO24121
C 7	99	10.4	754	12	BJ016176	BJ016176 BJO16176
C 8	98.5	10.3	961	10	BF203316	BF203316 601865914
C 9	98.5	10.3	1031	14	CB950999	CB950999 AGENCOURT
C 10	98.5	10.3	1141	11	AK080545	AK080545 MUS MUSC1
C 11	97.5	10.2	779	10	BF631437	BF631437 HVSMB001
C 12	96.5	10.1	844	11	CNS09D45	AX031096 Single re
C 13	96	10.1	701	10	BF863244	BF863244 963042C02
C 14	95	10.1	546	10	BF182274	BF182274 601804028
C 15	95.5	10.0	901	10	BF307233	BF307233 601891502
C 16	95.5	10.0	958	10	BG420860	BG420860 602452062
C 17	95	10.0	407	9	AW785806	AW785806 117260 MA
C 18	95	10.0	460	14	CB983286	CB983286 H001M02W
C 19	94.5	9.9	539	28	BH349665	BH349665 CH230-65E
C 20	94	9.9	582	14	CB286751	CB286751 CMD45_C08
C 21	94	9.9	1052	10	EG398041	BG398041 602439571
C 22	94	9.9	1263	13	BQ709745	BQ709745 AGENCOURT
C 23	93.5	9.8	736	12	BI768830	BI768830 603057734
C 24	93.5	9.8	938	13	BQ894657	BQ894657 AGENCOURT
C 25	93.5	9.8	953	9	AL555424	AL555424 AL555424
C 26	93	9.8	457	29	CNS02YOL	AL219990 Tetradon
C 27	93	9.8	470	13	BQ758584	BQ758584 Ema07_SO
C 28	93	9.8	1018	12	BQ054587	BQ054587 AGENCOURT
C 29	93	9.8	1291	10	BE622016	BE622016 601440668
C 30	92.5	9.7	655	14	CB868789	CB868789 H009014W
C 31	92.5	9.7	1008	12	BI755608	BI755608 603027112
C 32	92.5	9.7	1411	11	BC020343	BC020343 Homo sapi
C 33	92	9.7	756	14	CD348815	CD348815 UT-M-FYC-
C 34	92	9.7	832	10	BG387051	BG387051 602454749
C 35	92	9.7	871	10	BG178418	BG178418 602330206
C 36	92	9.7	898	29	CNS01VRS	ALL69466 Tetradon
C 37	92	9.7	963	10	BF794182	BF794182 602255566
C 38	92	9.7	1329	13	BQ960995	BQ960995 AGENCOURT
C 39	92	9.7	1640	10	BF180599	BF180599 601808704
C 40	91.5	9.6	422	14	CB763743	CB763743 AMGNNUC:S
C 41	91.5	9.6	539	10	BE757615	BE757615 212104 MA
C 42	91.5	9.6	641	9	AU127824	AU127824 AU127824
C 43	91.5	9.6	691	10	BB632604	BB632604 BB632604
C 44	91.5	9.6	701	14	CD262790	CD262790 PSMO19XF
C 45	91.5	9.6	844	12	B1198486	B1198486 602760491

ALIGNMENTS

RESULT 1
BF304699/c
LOCUS 601888252F1 NH_MGC_17 Homo sapiens cDNA clone IMAGE:4122276 '5',
DEFINITION 884 bp mRNA linear EST 21-NOV-2000
ACCESSION BF304699.1 GI:11251586
VERSION BF304699.1
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 984)


```

REFERENCE 1 (bases 1 to 643)
AUTHORS Kohara,Y., Shin-I,T., Kimura,T., Narita,T., Jindo,T. and Takeda,H.
TITLE Medaka EST Project in Takeda's lab
JOURNAL Unpublished
COMMENT Contact: Tadasu Shin-i
Center For Genetic Resource Information
National Institute of Genetics
1111 Yata, Mishima, Shizuoka 411-8540, Japan
Tel: 81-559-81-6856
Fax: 81-559-81-6855
Email: tshini@genes.nig.ac.jp.

FEATURES
    source
        1..643
            /organism="Oryzias latipes"
            /mol_type="mRNA"
            /strain="Hd-rR"
            /db_xref="taxon:8090"
            /clone="MF01SSA143012"
            /sex="mixture of female and male"
            /tissue_type="whole embryo"
            /dev_stage="segmentation stage 20 - 25"
            /clone_lib="MF01SSA cDNA"
            /clone_id="171 a 143 c 148 g 176 t"

BASE COUNT 171 a 143 c 148 g 176 t
ORIGIN
Alignment Scores:
Pred. No.: 11.7 Length: 643
Score: 99.00 Matches: 42
Percent Similarity: 33.77% Conservatve: 9
Best Local Similarity: 27.81% Mismatches: 50
Query Match: 10.39% Indels: 50
DB: 12 Gaps: 7

US-09-965-594-1 (1-182) x BJ024121 (1-643)
QY 27 TysAsnGlnValGluGlyGluValGlnIleValSerThrAlaAlaGlnThrPheLeuAla 46
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 242 AAAAATGAGTAAACACCAACACACATGACATGATGTCGGTTCACGGGCT 302
QY 47 ThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAlaGlyThrArgThrIleAla 66
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 302 -----TCGTGGAGAACCTATACAGTTCCTCTTACAGCAACGCCA 343
QY 67 SerProLys-----GlyProValIleGlnMetTyrThrAsnValAspLys 81
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 344 GCTCTGCGCGCGAGGAGCTCTCTGGCCAGCTGTG----- 379
QY 82 AspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrPro----- 97
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 380 -----ACTCGTGGAGAGAGAGAGCGTACCGCCGAGGCTGTAGG 418
QY 98 -----CysThrCysGlySerSerAspLeuTyrLeuValThrArg----- 110
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 419 CTGCAGGATGCGGATGTGGCTCTGCT-----TTGGTCTGCTCTCTCGTGGATCA 469
QY 111 -----HisAlaAspValIleProValAlaArgArgGlyAspSer 123
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 470 TCTTCTCACCCTGACCTTCCACATCCAGGTGTCGCCACCGCTGGTCTCAGCGGATCA 529
QY 124 ArgGlySerLeuLeuSerProArg-----ProIleSerTyrLeuLysGlySerSer 140
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 530 AGAGCGCGGACAGGCGAGTCGGGGTGAATCTCTCAGGACGCTCTCAGCGGATCA 589
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAla 151
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 590 GGAGGACCGACCTCGTCGACAGCTCTCTCTGCA 622

RESULT 7
BJ016176 754 bp mRNA linear EST 05-DEC-2001
LOCUS BJ016176 MF01SSA cDNA Oryzias latipes cDNA clone MF01SSA025C02 3'
DEFINITION mRNA sequence.
ACCESSION BJ016176

```

```

VERSION BJ016176.1 GI:17376695
KEYWORDS Oryzias latipes (Japanese medaka)
SOURCE Oryzias latipes
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Acanthomorpha; Acanthopterygii; Percormorpha; Atherinomorpha;
Belontiiformes; Adrianichthyidae; Oryziinae; Oryzias.
REFERENCE 1 (bases 1 to 754)
AUTHORS Kohara,Y., Shin-I,T., Kimura,T., Narita,T., Jindo,T. and Takeda,H.
TITLE Medaka EST Project in Takeda's lab
JOURNAL Unpublished
COMMENT Contact: Tadasu Shin-i
Center For Genetic Resource Information
National Institute of Genetics
1111 Yata, Mishima, Shizuoka 411-8540, Japan
Tel: 81-559-81-6856
Fax: 81-559-81-6855
Email: tshini@genes.nig.ac.jp.

FEATURES
    source
        1..754
            /organism="Oryzias latipes"
            /mol_type="mRNA"
            /strain="Hd-rR"
            /db_xref="taxon:8090"
            /clone="MF01SSA025C02"
            /sex="mixture of female and male"
            /tissue_type="whole embryo"
            /dev_stage="segmentation stage 20 - 25"
            /clone_lib="MF01SSA cDNA"
            /clone_id="194 a 151 c 181 g 198 t"

BASE COUNT 194 a 151 c 181 g 198 t
ORIGIN
Alignment Scores:
Pred. No.: 14.4 Length: 754
Score: 99.00 Matches: 42
Percent Similarity: 33.77% Conservatve: 9
Best Local Similarity: 27.81% Mismatches: 50
Query Match: 10.39% Indels: 50
DB: 12 Gaps: 7

US-09-965-594-1 (1-182) x BJ016176 (1-754)
QY 27 TysAsnGlnValGluGlyGluValGlnIleValSerThrAlaAlaGlnThrPheLeuAla 46
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 242 AAAAATGAGTAAACACCAACACACATGACATGATGTCGGTTCACGGGCT 302
QY 47 ThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAlaGlyThrArgThrIleAla 66
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 302 -----TGTGGAGAACCTATACAGTTCCTCTTACAGCAACGCCA 343
QY 67 SerProLys-----GlyProValIleGlnMetTyrThrAsnValAspLys 81
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 344 GCTCTGCGCGCGAGGAGCTCTCTGGCCAGCTGTG----- 379
QY 82 AspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrPro----- 97
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 380 -----ACTCGTGGAGAGAGAGAGCGTACCGCCGAGGCTGTAGG 418
QY 98 -----CysThrCysGlySerSerAspLeuTyrLeuValThrArg----- 110
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 419 CTGCAGGATGCGGATGTGGCTCTGCT-----TTGGTCTGCTCTCTCGTGGATCA 469
QY 111 -----HisAlaAspValIleProValAlaArgArgGlyAspSer 123
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 470 TCTTCTCACCCTGACCTTCCACATCCAGGTGTCGCCACCGCTGGTCTCAGCGGATGG 529
QY 124 ArgGlySerLeuLeuSerProArg-----ProIleSerTyrLeuLysGlySerSer 140
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 530 AGAGCGCGGACAGGCGAGTCGGGGTGAATCTCTCAGGACGCTCTCAGCGGATCA 589
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAla 151
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

```



```

Db      452 TG-----GGCACACTGTGCTCGTCCGACAT-----ATCGATCGCCCT 490
QY      69 LysGlyProValIleGlnMetIyrThrAsnValAspLysAsp.euValGlyTyrProAla 88
      ||||| ||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
Db      491 AAGAGCGCTTTCACAAAACACTT-----ACCTCCTTCCCTGCGCTGGC 535
QY      89 Pro-----GlnGlySerArgSerLeuThrProCysThrCysGly 101
      ||||| ||||| :||| :||| :||| :||| :||| :||| :||| :|||
Db      536 AT-GTTGGCAAAAAGAACCGCTTTTGGCTTCGCGCTTGGCGCCGCCCAATTGGGA 594
QY      102 SerSerAspLeuTyrLeuValThrArgHisAlaAsp-ValIleProValArgArgGcl 12:
      :||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
Db      595 ACCAGTGGC-----ACCACCATGGGCTTTGTGTGTCCTCCCTCCGCTGG 642
QY      121 YaspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerGcl 141
      :||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
Db      643 GCAATTACAAAACNCCCTTAACCGTCCGCTCGACACAAATTTCTTANGGCTCTGCA 702
QY      141 y-----GlyProLeuLeuLeuCysProAlaGlyHisAlaValGly 153
      ||||| ||||| :||| :||| :||| :||| :||| :||| :||| :|||
Db      703 TTTCCTTAAGTCCCGCTTTTSTTACCCACACACACATTTGTGCA 748

RESUL1 10
LOCUS   AK060545
DEFINITION Mus musculus 7 days neonate cerebellum cDNA, RIKEN full-length
          enriched library, clone:A730082L10 product:weakly similar to zinc
          finger protein (fragment) [Mus musculus], full insert sequence.
ACCESSION AK060545
VERSION   AK060545.1 GI:26348600
KEYWORDS  HTG; CAP trapper
SOURCE    Mus musculus (house mouse)
ORGANISM  Mus musculus
REFERENCE
AUTHORS   Carninci,P. and Hayashizaki,Y.
TITLE     High-efficiency full-length cDNA cloning
JOURNAL   Meth. Enzymol. 363, 19-44 (1999)
MEDLINE   99279253
PUBMED    10349536
REFERENCE
AUTHORS   Carninci,P., Shibata,Y., Hayashi,N., Sugahara,Y., Shibata,K.,
          Itoh,M., Konno,H., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
TITLE     Normalization and sub-traction of cap-trapper-selected cDNAs to
          prepare full-length cDNA libraries for rapid discovery of new genes
JOURNAL   Genome Res. 10 (10), 1517-1530 (2000)
MEDLINE   20499374
PUBMED    11042159
REFERENCE
AUTHORS   Shibata,K., Itoh,M., Aizawa,K., Nagaoka,S., Sasaki,N., Carninci,P.,
          Konno,H., Akiyama,J., Nishi,K., Katsunai,T., Tashiro,H., Itoh,M.,
          Sumi,N., Ishii,Y., Nakamura,S., Hazama,K., Nishino,I., Harada,A.,
          Yamamoto,K., Katsunoto,H., Sakaguchi,S., Ikegami,T., Kashiwagi,K.,
          Fujisawa,S., Inoue,K., Togawa,Y., Izawa,M., Obara,S., Watahiki,M.,
          Yoneda,Y., Ishikawa,T., Ozawa,K., Tanaka,T., Matsura,S., Kawai,S.,
          Okazaki,Y., Muramatsu,M., Inoue,Y., Kira,A. and Hayashizaki,Y.
TITLE     RIKEN integrated sequence analysis (RISA) system--384-format
          sequencing pipeline with 384 multicapillary sequencer
JOURNAL   Genome Res. 10 (11), 1757-1771 (2000)
MEDLINE   20530913
PUBMED    11076861
REFERENCE
AUTHORS   Kawai,J., Shinagawa,A., Shibata,K., Yoshino,M., Itoh,M., Ishii,Y.,
          Arakawa,T., Hara,A., Fukunishi,Y., Konno,H., Adachi,J., Fukuda,S.,
          Aizawa,K., Izawa,M., Nishi,K., Kiyosawa,H., Kondo,S., Yamanaka,I.,
          Saito,T., Okazaki,Y., Gojobori,T., Bono,H., Kasukawa,T., Saito,E.,
          Kadota,K., Matsuda,H., Ashburner,M., Batalov,S., Casavant,T.,
          Fleischmann,W., Gaasterland,T., Gissi,C., King,R., Kochiwa,H.,
          Kuehl,P., Lewis,S., Matsuo,Y., Nikaic,T., Pesole,G.,
          Quackenbush,J., Schram,I.M., Staubli,F., Suzuki,K., Tomita,M.,
          Wagner,L., Washio,T., Sakai,K., Okido,T., Furuno,M., Aono,H.,
          Baldarelli,R., Barsh,G., Blake,J., Boffelli,S., Bojunga,N.,

```

```

Carninci,P., de Bonaldo,M.F., Brownstein,M.J., Bult,C.,
Fletcher,C., Fujita,M., Gariboldi,M., Gustincich,S., Hill,D.,
Hofmann,M., Hume,D.A., Kamiya,M., Lee,N.H., Lyons,P.,
Marchionni,L., Mashima,J., Marzarelli,J., Mombaerts,P., Nordone,P.,
Ring,B., Ringwald,M., Rodriguez,I., Sakamoto,N., Sasaki,H.,
Sato,K., Schonbach,C., Seya,T., Shibata,Y., Storch,K.F., Suzuki,H.,
Toto-oka,K., Wang,K.H., Weitz,C., Whittaker,C., Wilming,L.,
Wynshaw-Boris,A., Yoshida,K., Hasegawa,Y., Kawaji,H., Kohtsuki,S.
and Hayashizaki,Y.
Functional annotation of a full-length mouse cDNA collection
Nature 409 (6821), 685-690 (2001)
21085660
1127851
5
The FANTOM Consortium and the RIKEN Genome Exploration Research
Group Phase 1 & II Team.
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
Nature 420, 563-573 (2002)
6 (bases 1 to 1141)
Adachi,J., Aizawa,K., Akimura,T., Arakawa,T., Bono,H., Carninci,P.,
Fukuda,S., Furuno,M., Hanagaki,T., Hara,A., Hashizume,W.,
Hayashida,K., Hayatsu,N., Hiramoto,K., Hiraoka,T., Hirozane,I.,
Hori,F., Imotani,K., Ishii,Y., Itoh,M., Kagawa,T., Kasukawa,T.,
Kato,H., Kawai,S., Kojima,Y., Kondo,S., Konno,H., Kouda,M.,
Koya,S., Kurihara,C., Matsuyama,T., Miyazaki,A., Murata,M.,
Nakamura,M., Nishi,K., Nomura,K., Numazaki,K., Ohno,M., Ohsato,N.,
Okazaki,Y., Saito,R., Saitoh,H., Sakai,C., Sakai,K., Sakazume,N.,
Sano,H., Sasaki,B., Shibata,K., Shinagawa,A., Shiraki,T.,
Sogabe,Y., Tagami,M., Tagawa,A., Takahashi,F., Takaku-Akahira,S.,
Takeda,Y., Tanaka,T., Tomaru,A., Toya,T., Yasunishi,A.,
Muramatsu,M. and Hayashizaki,Y.
Direct Submission
Submitted (16-APR-2002) Yoshihide Hayashizaki, The Institute of
Physical and Chemical Research (RIKEN), Laboratory for Genome
Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
RIKEN Yokohama Institute, 2-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.go.jp,
URL:http://genome.gsc.riken.go.jp/, Tel:81-45-503-9222,
Fax:81-45-503-9216)
CDNA library was prepared and sequenced in Mouse Genome
Encyclopedia Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in RIKEN.
Division of Experimental Animal Research in Riken contributed to
prepare mouse tissues.
Please visit our web site for further details.
URL:http://genome.gsc.riken.go.jp/
URL:http://fantom.gsc.riken.go.jp/
Location/Qualifiers
1..1141
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="FANTOM_DB:A730082L10"
/db_xref="taxon:10090"
/clone="A730082L10"
/tissue_type="cerebellum"
/clone_lib="RIKEN full-length enriched mouse cDNA library"
/dev_stage="7 days neonate"
<1..587
/note="unnamed protein product; putative
weakly similar to zinc finger protein (fragment) [Mus
musculus] (PIR|I48722, evidence: FASTY, 50.7%ID,
57.6%length, match=601)"
/codon_start=3
/protein_id="BAC37940.1"
/db_xref="GI:26348601"
/translation="DSCLPASGRSLTTPRGDGFLEKLSAARAVGPGSPVAFGV
TRYGAQAGQRRRCACRRSGCLCLSRKPRRHVPVPGVHYGLSGRRIPPPAGE
AQAGRAPQOVPHPGCRPHGTVVPGGAAGLLPALAAROVPGVPGREGPRAPRHS
KRPVPTALGFSGCGGAPAPLLAPANGRSVGLAL"
1118..1123
/note="putative"
polyA_signal

```


DNA Sequencing By: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov
 Plate: LLCMI044 row: c column: 02
 High quality sequence start: 6
 High quality sequence stop: 684.

FEATURES

Location/Qualifiers
 1..901
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:4137145"
 /tissue_type="rhabdomyosarcoma"
 /lab_host="p810B (phage-resistant)"
 /clone_lib="NIH_MGC_13"

/note="Organ: muscle; Vector: pOTB7; Site_1: EcoRI;
 Site_2: XhoI; cDNA made by oligo-dT priming.
 Directionally cloned into EcoRI/XhoI sites using the
 following 5' adaptor: GGCACGAG(G). Size-selected >500bp
 for average insert size 1.8kb. Library constructed by
 Ling Hong in the laboratory of Gerald M. Rubin (University
 of California, Berkeley) using ZAP-cDNA synthesis kit
 (Stratagene) and Superscript II RT (Life Technologies)."

BASE COUNT 144 a 267 c 329 q 161 t

ORIGIN

Alignment Scores:
 Pred. No.: 39.5 Length: 901
 Score: 95.50 Matches: 37
 Percent Similarity: 38.46% Conservative: 8
 Best Local Similarity: 31.62% Mismatches: 28
 Query Match: 10.02% Indels: 45
 DB: 10 Gaps: 5

US-09-965-594-1 (1-182) x BF307233 (1-901)

QY 52 ValCysTrpIhrValIyrHis-----GlyAlaGlyThrArgThrIleAl 66
 Db 620 ATGTGCTGGACGGGTCCCGCCGCGCATCTCTAGCGGGGGTCCGGCAGCCAGCGGTGG 679
 QY 66 aSerProLysGlyProValIleGlnMetTyrThrAsnValAspLysAspLeuValGlyTr 86
 Db 680 TGGGCAGGACGGTGGAGTGTGCA-----GTCACAGGATG 715
 QY 86 pProAlaProGlnGlySerArgSerLeuThrProCysThrCysGlySerAspLeuTy 106
 Db 716 GCCCGCCCATCCGG----- 731
 QY 106 rLeuValIhrArgHisAlaAspValIleProValArgAlaGlyAspSerArgGlySe 126
 Db 732 -----GTGAGCCTTCCTCAGGGGGTTCGGGGGGGTTC 756
 QY 126 rLeuLeuSerProArgProIleSerTyrIleLysGlySerSerGlyGlyProIleLeuCy 146
 Db 767 CATTTCGGTCCGA-----GGGGAATCTCTCCAGGGGCGCTGGACTG 810
 QY 146 sProAlaGlyHisAlaValGly-----IlePheArgAlaAlaValCys 160
 Db 811 TCCGGCGGGTCCCGCGCGCCGACACAGCGGTCCGGCGGCTGTGC 859

Search completed: September 27, 2003, 14:09:39
 Job time : 2238 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - nucleic search, using frame_plus_p2n mode:

Run on: September 27, 2003, 12:22:20 : Search time 327 Seconds
(without alignments)
1502.44c Million cell. updates/sec

Title: US-09-965-594-1

Perfect score: 953

Sequence: 1 MAPITAYACGTCRLGCIIT.....GVAKAVCFIPVSELETINKS 182

Scoring table:

MIOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 2552756 seqs, 1349719017 residues

Total number of hits satisfying chosen parameters: 5105511

Minimum DB seq length: 9

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 99%

Listing first 45 summaries

Command line parameters:

-MODEL=frame+ p2n.model -DEV=xlh
-Q/cgn2_1/USPTO.spool/US0965594/runat_26092003_164342_18746/app_query.fasta_1.327
-DB=N_Geneseq_19Jun03 -QFMT=fastap -SUFFIX=ring -MINMATCH=0.1 -LOOPCL=0
-LOOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=bloms62 -TRANS=human40.cdi
-LIST=45 -DOCALIGN=200 -THR SCORE=pc1 -THR MAX=99.9 -THR MIN=0 -ALIGN=15
-MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEAPS=2E+500 -MINLEN=0 -MAXLEN=200000000
-USER=US0965594 -CGC=1.1.2 -runat_26092003_164342_18746 -NPU=6 -ICP=1
-NO_MMAPP -LARGEQUERY -NEG_SCORES=0 -WAIT -DSPLOC=100 -LONGLOG
-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database : N_Geneseq_19Jun03.*

1: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1980.DAT.*
2: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1981.DAT.*
3: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1982.DAT.*
4: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1983.DAT.*
5: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1984.DAT.*
6: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1985.DAT.*
7: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1986.DAT.*
8: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1987.DAT.*
9: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1988.DAT.*
10: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1989.DAT.*
11: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1990.DAT.*
12: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1991.DAT.*
13: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1992.DAT.*
14: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1993.DAT.*
15: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1994.DAT.*
16: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1995.DAT.*
17: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1996.DAT.*
18: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1997.DAT.*
19: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1998.DAT.*
20: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA1999.DAT.*
21: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA2000.DAT.*
22: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
23: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
24: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA2002.DAT.*
25: /SIDSL/qcgdata/geneseq/geneseq-emb1/NA2003.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed.

and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
1	946	99.3	2058	24	AAD29795	HCV-1 NS3/4a mutan
2	946	99.3	2058	24	ABK15344	Hepatitis C virus
3	946	99.3	2058	25	ABX14410	DNA encoding HCV-1
4	943	99.0	5300	10	AAAN92097	Combined open read
5	943	99.0	5360	10	AAAN90327	Hepatitis C virus
6	943	99.0	6905	10	AAAN92103	Combined open read
7	943	99.0	7310	10	AAAN92106	Combined open read
8	943	99.0	7310	10	AAAN90336	Composite hepatitis
9	943	99.0	7310	16	AAQ98221	Hepatitis C virus
10	943	99.0	8316	21	AAA75296	cDNA sequence comp
11	943	99.0	9133	20	AAZ07656	Nucleotide sequenc
12	943	99.0	9185	11	AAQ05956	Sense strand of th
13	943	99.0	9185	12	AAQ10566	Hepatitis C virus
14	943	99.0	9185	21	AAA75297	Sense strand of HC
15	943	99.0	9400	13	AAQ21744	Compiled HCV cDNA.
16	943	99.0	9401	17	AAT12710	Hepatitis C virus
17	943	99.0	9401	18	AAT99981	HCV polyprotein co
18	943	99.0	9401	19	AAV09989	HCV polyprotein co
19	943	99.0	9401	24	AAD35043	Hepatitis C virus
20	942	98.8	2061	24	AAD34500	Hepatitis C virus
21	942	98.8	2061	24	AAD31767	Hepatitis C virus
22	940	98.6	9185	20	AAZ26737	Nucleotide sequenc
23	940	98.6	9185	20	AAZ00459	Hepatitis C virus
24	939	98.5	8316	11	AAQ05955	Hepatitis C virus
25	939	98.5	9502	15	AAQ74770	Hepatitis C virus
26	937	98.3	9646	19	AAV59361	Hepatitis C virus
27	937	98.3	9646	24	ABK87285	cDNA encoding hepa
28	937	98.3	12380	19	AAV59364	Hepatitis C virus
29	937	98.3	12980	24	ABK87286	Hepatitis C virus
30	937	98.3	16522	21	AAZ36212	Nucleotide sequenc
31	936	98.2	630	17	AAI43708	Plasmid pT5H18/HIV
32	936	98.2	630	17	AAI43294	HCV insoluble NS3
33	936	98.2	630	18	AAI58401	HCV NS3 protease c
34	936	98.2	810	17	AAT38903	pNH8deltaA H7 e
35	936	98.2	810	17	AAT42389	HCV soluble NS3 pr
36	936	98.2	810	18	AAT58396	HCV NS3A ORF comp
37	935	98.1	6299	22	AAE83669	HCV NS3 DNA
38	934	98.0	1933	20	AAZ23258	Plasmid pMT-BS(+)/
39	934	98.0	8145	20	AAZ23259	Hepatitis C virus
40	933	97.9	594	21	AAA73335	Hepatitis C virus
41	933	97.9	9365	24	AAZ25518	Hepatitis C virus
42	933	97.9	9401	17	AAT41882	Hepatitis C virus
43	933	97.9	9416	19	AAV59378	Hepatitis C virus
44	933	97.9	9416	24	ABK87300	cDNA encoding hepa
45	931	97.7	549	17	AAT43706	Hepatitis C virus

ALIGNMENTS

RESULT 1
AAD29795 AAD29795 standard; DNA; 2058 BP.
XX
AC AAD29795;
XX
DT 17-MAY-2002 (first entry)
XX
CE HCV-1 NS3/4a mutant conformational antigen encoding DNA.
XX
KW Hepatitis C virus; NS3/4a antigen; HCV infection; mutant; ds.
XX
OS Hepatitis C virus type 1.
XX
FH Synthetic.
FT CDS
Key Location/Qualifiers
1..686

```

/*tag= a
/product= "HCV-1 NS3/4a conformational antigen"
/Note= "CDS does not include stop codon"
/partial

WO200196875-A2.
20-DEC-2001.

14-JUN-2001: 2001WO-US19169.
15-JUN-2000: 2000US-212082P.
02-APR-2001: 2001US-280811P.
02-APR-2001: 2001US-280867P.

(CHIR ) CHIRON CORP.

Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Coit D;
Medina-Selby A;
WPI: 2002-179522/23.
P-PSDB; AAE18689.

Immunosay solid support useful for detecting hepatitis C virus
infection in a biological sample, comprises at least one of HCV
anti-core antibody and HCV NS3/4a epitope, bound to the support .

Example 2: Fig 4; 87pp; English.

The present invention relates to hepatitis C virus (HCV) core antigen
and NS (nonstructural) 3/4a antibody combination assay that can detect
both HCV antigens and antibodies present in a sample using a single
solid matrix as well as immunosay solid supports for use in the assay.
The solid support is useful for detecting HCV infection in a biological
sample. The present sequence is a DNA encoding HCV-1 NS3/4a mutant
conformational antigen. This sequence is used in the exemplification
of the invention.

SQ Sequence 2058 BP: 419 A: 634 C: 590 G: 425 T: 0 other:

Alignment Scores:
Pred. No.: 1,468-78 Length: 2058
Score: 945.00 Matches: 190
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 98.90% Mismatches: 0
Query Match: 95.27% Indels: 0
DB: 24 Gaps: 0

us-09-965-594-1 (1-182) x AAD29795 (1-2058)

QY 1 MetAlaProIleThrAlaTyrAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
DB 1 ATFGCGCCCATCACGGCGTAGCCCGAGAGCAAGGGGCCCTCTAGGGTGCATATCACC 60
QY 21 SerLeuThrGlyArgAspGlyAsnGlnValGlnGlyGluValGlnIleValSerThrAla 40
DB 61 AGCCTAACTGGCGCGGACAAACCAAGTGGAGGTGAGTCCAGATTGTGCTCAACTGCT 120
QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpIleValIleHisLeuAla 60
DB 121 GCCCAAACTTCTCGACACGTGCATCAATAGGGTGTCTGACGTGTACACCGGGGCC 180
QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
DB 181 GGACGAGGACCATCGCTCACCCAGGGTCTGTGATCCAGATGTATACCAATGTAGAC 240
QY 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB 241 CAAGACCTGTGGGTGGCGCCGTCGCAAGTACCGCATTCATGCACCTGCACTTCC 300
QY 101 GlySerSerAspLeuTyrIleuValThrArgHsAlaAspValIleProValArgArgArg 120
DB 301 GGCTCTCGGACCTTACCTGTGCAGAGGACGCGCATGTCTCCGTGCGGCGCG 360

QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB 361 GGTGATAGCAGGGGCGAGCCTGTGTGCCCCCGGCCCATTTCCCTACTGAAAGGCTCCICG 420
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValIleGlyPheArgAlaAlaValCys 160
DB 421 GGGGGTCCGCTGTGTGTGCCCCGGGCGACCGCGTGGCATATTTAGGGCGCGGTGTGC 480
QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValGluSerLeuGluThrMet 180
DB 481 ACCCGTGGAGTGGCTAAGCGGTGGACTTATCCCTGTGGAGAACCTAGAGACACCATG 540
QY 181 ArgSer 182
DB 541 AGGTCC 546

RESULT 2
ABK15344
1C ABK15344 standard; DNA: 2058 BP.
XX
AC ABK15344:
XX
DE 08-MAY-2002 (first entry)
DE Hepatitis C virus NS3/4a conformational epitope gene sequence.
XX
KW Hepatitis C virus; HCV; NS3/4a conformational epitope; seroconversion;
KW immunosay solid support; multiple epitope fusion antigen; MEFA;
KW non-structural protein; gene; ds.
XX
OS Hepatitis C virus.
XX
FH Key Location/Qualifiers
ET CDS 1..2058
FI /tag= a
FI /product= "HCV NS3/4a conformational epitope"
FI /note= "This sequence lacks a stop codon"
XX
PN WO200196870-A2.
XX
PD 20-DEC-2001.
XX
PF 14-JUN-2001; 2001WO-US19156.
XX
PR 15-JUN-2000; 2000US-212082P.
PR 02-APR-2001; 2001US-280811P.
PR 02-APR-2001; 2001US-280867P.
XX
PA (CHIR ) CHIRON CORP.
XX
PI Chien DY, Arcangel P, Tandeske L, George-nascimento C, Coit D;
PI Medina-selby A;
PI
XX
WP: 2002-090228/12.
DR P-PSDB; AAU76377.
XX
PT Immunosay solid support, useful for detecting hepatitis C virus
infection in biological sample, comprises HCV NS3/4a conformational
epitope and multiple epitope fusion antigen bound to the support -
XX
PS Disclosure: Fig 3; 92pp; English.
XX
CC The present invention relates to a new immunosay solid support
consisting essentially of at least one hepatitis C virus (HCV) NS3/4a
conformational epitope and a multiple epitope fusion antigen (MEFA),
bound to the support. The NS3/4a conformational epitope and/or
MEFA reacts specifically with anti-HCV antibodies present in a biological
sample from an HCV-infected individual. The immunoassay of the invention
is useful for detecting hepatitis C virus infection in a biological
sample. The method of the invention provides a sensitive, accurate
diagnostic and prognostic tool to provide adequate patient care and to

```

CC prevent transmission of HCV by blood and by blood products, or by
 CC personal contact. Use of NS3/4a conformational epitope in combination
 CC with MEFA, provides a sensitive and reliable method for detecting early
 CC HCV seroconversion. Use of MEFA has the added advantages of decreasing
 CC masking problems, improving sensitivity in detecting antibodies by
 CC allowing a greater number of epitopes on a unit surface area of
 CC substrate, and improving substrate. Detection accuracy is increased and
 CC the incidence of false results is reduced because of the identification
 CC and use of highly immunogenic HCV antigens which are present during
 CC the early stages of HCV seroconversion. The present nucleic acid sequence
 CC encodes the non-structural protein NS3/4a conformational epitope of the
 CC invention.
 XX
 SQ Sequence 2058 BP; 419 A; 633 C; 581 G; 425 T; 0 other;

Alignment Scores:
 Pred. No.: 1,46e-78 Length: 2058
 Score: 946.00 Matches: 180
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 98.90% Mismatches: 0
 Query Match: 99.27% Indels: 0
 DB: 24 Gaps: 0

US-09-965-594-1 (1-182) x ABX15344 (1-2058);

QY 1 MetAlaProIleThrAlaValAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
 Db 1 ATGGCCCATCATCGGCGTACGCGCCAGCAGCAAGGGGCTCTCTAGGTGCTAATCAC 50
 QY 21 SerLeuThrGlyArgAspLysAsnGlnValGlnGlyValGlnIleValSerThrAla 40
 Db 51 AGCTTAACGGCGGCGACAAACCAAGTGGAGGTGAGGTTCAGATGTGCACTGCT 120
 QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValIleHisGlyAla 60
 Db 121 GCCCAACCTCTCTGCAACGTCATCAATGGGTGTCTGCTGCTGCTACCAACGGG 180
 QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetThrThrAsnValAsp 80
 Db 181 GGACACGAGGACCATCGGTCGCCAGGAGGTCTGTCTATCATCAGATGATACCAATG 240
 QY 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
 Db 241 CAAGACCTTGCGGCTGGCGGCTGCGCAAGGTAGCGGATCATTCACACCTGCACT 300
 QY 101 GlySerSerAspLeuThrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
 Db 301 GGCTCTCCGAGACCTTACCTGGTCAGAGGACCGGATGTCATTCGGTGGCGCGGCG 360
 QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
 Db 361 GGTGATAGCAGGGGACGCTGCTGTCGCCCGCCGCTTCTACTTGAAGGCTCTCTG 420
 QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaIleValCys 160
 Db 421 GGGGTGCGGTGTGTGTGCGCGGGGCGGCGGCGGTGGGATATTAGCGCGCGGTGTC 480
 QY 161 ThrArgGlyValAlaCysAlaValAspPheIleProValGlnSerLeuLeuIleThrMet 180
 Db 481 ACCGTGGAGTGGCTAAGCGGTGGACATTATCCCTGTGAGAACCTAGACACCACTG 540
 QY 181 ArgSer 182
 Db 541 AGGTCC 546

RESULT 3

ID ABX14410 standard; DNA: 2058 BP.

XX ABX14410;

AC ABX14410;

XX 06-MAR-2003 (first entry)

DT

XX

DE DNA encoding HCV-1 NS3/4a conformational antigen.
 XX
 KW Immunoassay solid support; Hepatitis C Virus type-1; HCV-1;
 KW NS3/4a conformational epitope; multiple epitope fusion antigen;
 KW MEFA; anti-HCV antibody; NS3/4a conformational antigen;
 KW HCV infection; mutant; gene; ds.
 XX
 OS Hepatitis C virus type 1.
 OS Synthetic.
 FH
 FT Key location/Qualifiers
 FT CDS 1..2058
 FT /*tag= a
 FT /partial= "NS3/4a conformational antigen"
 FT /product= "NS3/4a conformational antigen"
 FT /note= "This sequence lacks a stop codon"
 XX
 US02002146685-A1.
 XX
 10-OCT-2002.
 XX
 14-JUN-2001; 2001US-0881654.
 XX
 15-JUN-2000; 2000US-212082P.
 PR 02-APR-2001; 2001US-280811P.
 PR 02-APR-2001; 2001US-280867P.
 XX
 (CHIE/) CHIEN D Y.
 PA (ARCA/) ARCANDEL P.
 PA (TAND/) TANDESKE L.
 PA (GEOR/) GEORGE-NASCIMENTO C.
 PA (COIT/) COIT D.
 PA (MEDI/) MEDINA-SELBY A.
 XX
 PI Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Coit D;
 Medina-Selby A;
 DR WP1: 2003-147573/14.
 DR P-PSDB; ABG72261.
 XX
 PI Immunoassay solid support for detecting Hepatitis C Virus infection in
 PI biological samples, comprises Hepatitis C Virus conformational epitope
 PI and multiple epitope fusion antigen -
 XX
 PS Disclosure: Fig 3A-3D; 45pp; English.
 XX
 CC The present invention relates to immunoassays comprising Hepatitis C
 CC Virus (HCV) NS3/4a conformational epitope and multiple epitope fusion
 CC antigen (MEFA), bound to a solid support. The NS3/4a epitope and/or
 CC the multiple epitope fusion antigen react with anti-HCV antibodies
 CC present in a biological sample from an HCV-infected individual. The
 CC immunoassays and methods of the invention are useful for detecting
 CC HCV infection in a biological sample. The inventive immunoassay solid
 CC support provides a sensitive and reliable method for detecting early
 CC HCV seroconversion. The assays can detect HCV infection caused by any
 CC six known genotypes of HCV. The use of the multiple epitope fusion
 CC proteins decreases masking problems, improves sensitivity in detecting
 CC antibodies by allowing a greater number of epitopes on a unit area
 CC of substrate, and improves selectivity. The present sequence
 CC encodes HCV type 1 (HCV-1) NS3/4a conformational antigen, a mutant of
 CC the HCV-1 NS3/4a polypeptide.
 XX
 SQ Sequence 2058 BP; 419 A; 633 C; 581 G; 425 T; 0 other;

Alignment Scores:

Pred. No.: 1,46e-78 Length: 2058
 Score: 946.00 Matches: 180
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 98.90% Mismatches: 0
 Query Match: 99.27% Indels: 0
 DB: 25 Gaps: 0

US-09-965-594-1 (1-182) x ABX14410 (1-2058)

Qy 1 MetAlaProIleThrAlaTyrAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
 Db ATGCGGCCCATCACGGCTACGGCCAGCAGACAGAGGGGCTCTAGGTGCATAATCACC 60
 Qy 21 SerLeuThrGlyArgAspLysAsnGlnValGlnGlyValGlnIleValSerThrAla 40
 Db AGCCTAACTGGCGGGGCAAAACCAAGTAGTGAGGGTGCAGATGCTGCAACTGCT 120
 Qy 41 AlaGlnThrPheLeuAlaThrCysIleAspGlyValCysIleThrValTyrHisGlyAla 60
 Db GCCCAACCTTCCTGGCAACCTGCATCAATGGGCTGTCTGGACTGCTACAGGGGGCC 180
 Qy 61 GlyThrArgThrIleAlaSerProLysGlyProValIleSerMetTyrThrAsnValAsp 80
 Db GGAACGAGGACCATCGGCTCACCAAGGGTCTCTCATCCAGATGATACCAATGTAGAC 240
 Qy 81 LysAspLeuValGlyTyrProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
 Db CAAGACCTTGGGCTGGCGCGCTCCCGCAAGTAGCCGATCATTGACACCCGCACTTGC 300
 Qy 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
 Db GGCTCCCGGACCTTTACCTGGTCACGAGCAGCCGCACTCATTCGCGCGCGCGCG 360
 Qy 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuGlyGlySerSer 140
 Db GGTGATACGAGGCGACGCTGCTGTCCCGCGCGCCCATCTCTACITGAAAGGCTCTCG 420
 Qy 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
 Db GGGGTCGCGCTGTGTGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 480
 Qy 161 ThrArgGlyValAlaLysAlaValAspPheIleProValGluSerLeuGluThrThrMet 180
 Db ACCGCTGGAGTGGCTAAGCGGTGGACATTATCCCTGTGGAGACCTAGAGACACCATG 540
 Qy 181 ArgSer 182
 Db AGGTCC 546

RESULT 4
 AAN92097
 ID AAN92097 standard: DNA: 5300 BP.
 XX AAN92097;
 XX 25-MAR-2003 (updated)
 DI 02-MAR-1990 (first entry)
 DE Combined open reading frames of the hepatitis C virus (HCV) cDNA in
 DE clones 14i, 11b, 7f, 7e, 8h, 33c, 40b, 37b, 35, 36, 81, 32, 33b, 25c, 14c.
 DE 8f, 33f, 33g and 39c.
 XX Hepatitis C virus: HCV; non-A, non-B hepatitis; NANBH.
 XX Hepatitis C virus.
 OS
 FH Key Location/Qualifiers
 CDS 3..5300
 FT /tag- a
 XX
 PN EP318216-A.
 XX
 PD 31-MAY-1989.
 XX
 PF 18-NOV-1988; 88EP-0310922.
 XX
 PR 16-NOV-1987; 87US-0122714.
 PR 30-DEC-1987; 87US-0139886.
 PR 26-FEB-1988; 88US-0161072.
 PR 06-MAY-1988; 88US-0191251.
 PR 26-OCT-1988; 88US-0263584.

PR 14-NOV-1988; 88US-0271450.
 XX
 PA {CHIR } CHIRON CORP.
 XX
 PI Houghton M, Choo QL, Kuo G;
 XX
 XX MPI; 1989-159274/22.
 DR P-PSDB; AAP92041.
 DR
 XX
 PT Purified hepatitis C virus
 PT - and associated nucleic acids and polypeptide(s)
 PT

PS Claim 3: Figure 26-1, 26-2, 26-3, 26-4, 26-5, 26-6; 139pp: English.

XX It is a double-stranded nucleotide sequence of the open reading frame (ORF) (tag a) extending through clones 14i, 11b, 7f, 7e, 8h, 33c, 40b, 37b, 35, 36, 81, 32, 33b, 25c, 14c, 8f, 33f, 33g and 39c of hepatitis C virus (HCV) cDNA. In creating the composite sequence the following heterogeneities were considered. Clone 33c contains a sequence of 800 base pairs which overlaps the cDNAs in clones 40b and 37c. In clone 33c, as well as in 5 other overlapping clones, nucleotide #789 is a G. However, in clone 37b the corresponding nucleotide is an A. This heterogeneity may have important ramifications for protein folding. Nucleotide #2 in clone 8h is a T which may represent a cloning artifact because the corresponding residue in clone 7e and in 3 other overlapping clones is an A. Therefore the residue in this position is designated as an A. The 3'-terminal nucleotide in clone 8f is represented as a T than a G because the corresponding residue in clone 33f and in 2 other overlapping clones is a T. The 3' terminal sequence of clone 33f is represented as ATTC, as is found in the corresponding sequence in clone 33g and in 2 other overlapping clones, rather than as TTGC, as is found in clone 33f. Residue #4 in clone 33g is designated as A rather than a T because the corresponding residue in clone 33f and 2 other overlapping clones is an A. The 3'-terminus of clone 14i is depicted as TA rather than AA because the corresponding dinucleotide in clone 11b and 3 other clones is TA. Potential cloning artifacts have been omitted and instead the corresponding sequences in non-5'-terminal regions of multiple overlapping clones are shown. AAN92097 could be used as a source of oligomeric DNA hybridisation probes to detect the presence of HCV nucleic acids in samples. The polypeptide(s) it encodes could be used as immuno- assay reagents and vaccines and to generate antibodies useful in diagnosis and passive immunotherapy for HCV infection/non-A, non-B hepatitis.
 CC (Updated on 25-MAR-2003 to correct PR field.)
 CC (Updated on 25-MAR-2003 to correct PI field.)
 XX

SQ Sequence 5300 BP; 1047 A; 1606 C; 1515 G; 1130 T; 2 other;

Alignment Scores:

Prod. No.: 8-9e-78 Length: 5300
 Score: 943.00 Matches: 179
 Percent Similarity: 100.00% Conservative: 3
 Best Local Similarity: 98.35% Mismatches: 0
 Query Match: 98.95% Indels: 0
 DB: 10 Gaps: 0

CS-09-965-594-1 (1-162) x AAN92097 (1-5300)

Qy 1 MetAlaProIleThrAlaTyrAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
 Db CTGGCGGCCCATCACGGCTACGGCCAGCAGACAGAGGGGCTCTAGGTGCATAATCACC 989
 Qy 21 SerLeuThrGlyArgAspLysAsnGlnValGlnGlyValGlnIleValSerThrAla 40
 Db AGCCTAACTGGCGGGGCAAAACCAAGTAGTGAGGGTGCAGATGCTGCAACTGCT 1049
 Qy 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysIleThrValTyrHisGlyAla 60
 Db GCCCAACCTTCCTGGCAACCTGCATCAATGGGCTGTCTGGACTGCTACAGGGGGCC 1109
 Qy 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
 Db GGAACGAGGACCATCGGCTCACCAAGGGTCTCTCATCCAGATGTATACCAATGTAGAC 1169

QY 81 LysAspLeuValaIyLrPpProlaProGingLySerArgSerLeuThrProCysThrCys 100
 ::
 Db 1170 CAAGACCTGTGGGCTGGCCGCTCGCAAGGTAGCGGTCAATTGACACCTGCACTTGC 1229
 ::
 QY 101 GlySerSerAspLeuTyLerValThrArgHisAlaAspValIleProValArgArg 120
 ::
 Db 1230 GGCTCTCGGACCTTTACCTGGTTCAGAGGACGCGGATGTGATTCCTCGCGCGGGCG 1269
 ::
 QY 121 GlyAspSerArgGlySerLeuSerProArgProIleSerTyLerCysGlySerSer 140
 ::
 Db 1290 GTGTAAGCAGGGCAGCGTGTGTGGCCCGGCGCAATTCCTACTTGAAGGCTCTGCG 1349
 ::
 QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyLePheArgAlaAlaValCys 160
 ::
 Db 1350 GGGGGTCCGCTGTGTGGCCCGCGGGCGACGGCTGGGCTATATTAGGCGCTGGGTGC 1409
 ::
 QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValGlySerLeuThrPro 180
 ::
 Db 1410 ACCCGTGGAGTGGCTTAGGCGGTGGACATTTCCTGTGGAGACCTAGAGACACCATG 1469
 ::
 QY 181 ArgSer 182
 ::::::::::
 Db 1470 AGGTCC 1475

RESULT 5
 AAN90327
 ID AAN90327 standard; cDNA; 5360 BP.
 AC AAN90327;
 DT 25-MAR-2003 (updated)
 DT 11-NOV-1989 (first entry)
 XX Hepatitis C virus composite probe.
 XX Hepatitis C virus: composite cDNA; probe; vaccine.
 XX Pan troglodytes.
 XX Key Location/Qualifiers
 FT CDS 3..5360
 FT /*tag= a
 XX GB22-2511-A.
 XX 26-JUL-1989.
 XX 18-NOV-1988; 88GB-0027024.
 XX 18-NOV-1987; 87US-0122714.
 XX 30-DEC-1987; 87US-0139886.
 XX 26-FEB-1988; 88US-0161072.
 XX 26-OCT-1988; 88US-0263584.
 XX (CHIR) CHIRON CORPORATION.
 XX Houghton M, Choo QL, Kuo G;
 XX WFI; 1989-215054/40.
 XX Hepatitis C virus gene - used for prodn. of polynucleotide probes,
 PT polypeptide(s) and antibodies for diagnosis, prevention and treatment
 PT of infection.
 XX Disclosure: Fig. 26; 174pp; English.
 XX The sequence shows the composite cDNA sequence derived from the aligned
 CC hepatitis C virus (HCV) cDNA's in clones 141, 11b, 7f, 8f, 33c, 40b,
 CC 37b, 35, 36, 8f, 32, 33b, 25c, 14c, 8f, 33f, 33g and 39c. The cDNA
 CC encodes antigens which react with antibodies in patients with non-A
 CC non-B hepatitis (NANBH). The cDNA can be used to design probes, or to
 CC synthesise polypeptides, which are used to diagnose HCV-induced NANBH,

CC to raise antibodies for immunoassay or treatment, or to produce
 CC vaccines. See also AAP90158, AAN90303-26, and AAN90328-36.
 CC (Updated on 25-MAR-2003 to correct PR field.)
 XX
 SQ Sequence 5360 BP; 1060 A; 1622 C; 1532 G; 1145 T; 1 other:
 Alignment Scores:
 Pred. No.: 9, 04e-78 Length: 5360
 Score: 943.00 Matches: 179
 Percent Similarity: 100.00% Conservative: 3
 Best Local Similarity: 98.35% Mismatches: 0
 Query Match: 98.95% Indels: 0
 Gaps: 10
 US-09-965-594-1 (1-182) x AAN90327 (1-5360)
 QY 1 MetAlaProIleThrAlaValAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
 ::
 Db 930 CTGGGCCCATCAGGGCTAGCCAGCAGACAGAGGGGCTCTCTAGGGTGCATATCACC 989
 ::
 QY 21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGluValGlnIleValSerThrAla 40
 ::
 Db 990 AGCCTAACTGGCGGGACAAAACCAAGTGCAGGCTAGGTCCAGANTGTGTCAACTGCT 1049
 ::
 QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValIleHisCysAla 60
 ::
 Db 1050 GCCCAAACTTCCTGGCAACGTGCATCAATGGGTGTGCTGGACGTGTACACGGGGGCC 1109
 ::
 QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyThrAsnValAsp 80
 ::
 Db 1110 GGNAGGAGGACCATCGCGTCACCAAGGCTCTGTCTATCCAGATGTATACCAATGTAGAC 1169
 ::
 QY 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
 ::
 Db 1170 CAAGACCTGTGGGCTGGCCGCTCGCAAGGTAGCGGTCAATTGACACCTGCACTTGC 1229
 ::
 QY 101 GlySerSerAspLeuTyLerValThrArgHisAlaAspValIleProValArgArg 120
 ::
 Db 1230 GGCTCTCGGACCTTTACCTGGTTCAGAGGACGCGGATGTGATTCCTCGCGGGCGG 1289
 ::
 QY 121 GlyAspSerArgGlySerLeuSerProArgProIleSerTyLerCysGlySerSer 140
 ::
 Db 1290 GTGTAAGCAGGGCAGCGTGTGTGGCCCGGCGCAATTCCTACTTGAAGGCTCTGCG 1349
 ::
 QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyLePheArgAlaAlaValCys 160
 ::
 Db 1350 GGGGGTCCGCTGTGTGGCCCGCGGGCGACGGCTGGGCTATATTAGGCGCGGGGTGC 1409
 ::
 QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValGlySerLeuThrPro 180
 ::
 Db 1410 ACCCGTGGAGTGGCTTAGGCGGTGGACATTTCCTGTGGAGACCTAGAGACACCATG 1469
 ::
 QY 181 ArgSer 182
 ::::::::::
 Db 1470 AGGTCC 1475

RESULT 6
 AAN92103
 ID AAN92103 standard; DNA; 6905 BP.
 XX AAN92103;
 AC AAN92103;
 XX 25-MAR-2003 (updated)
 DT 02-MAR-1990 (first entry)
 XX Combined open reading frames of the hepatitis C virus (HCV) cDNAs from
 DE clones 12f through 15e.
 XX Hepatitis C virus; HCV; non-A, non-B hepatitis; NANBH.
 XX Hepatitis C virus.
 XX Key Location/Qualifiers


```

Alignment Scores:
Pred. No.: 1,32e-77 Length: 7310
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservatives: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 10 Gaps: 0

US-09-965-594-1 (1-182) x AAN921106 (1-7310)
QY 1 MetAlaProIleThrAlaTyrAlaGlnInrArqGlyLeuLeuGlyCysLeuIleThr 20
DB 1728 CTGGCCGCCATCAGCGGTACGCCACAGACAGAGCGGCTCTAGGTGCATATCAC 1787
QY 21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyValGlnIleValSerThrAla 40
DB 1788 AGCCTAACTGGCGGACAAAACCAAGTGGAGGGTCAGTTCAGATTTGTGCAACTGCT 1847
QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAla 60
DB 1848 GCCCAAACTTCTGGCAACGTGCATCAATGGGG:GTGGTGGACTGTCTACCAACGGGGG 1907
QY 61 GlyThrArqThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
DB 1908 GGAACGAGGACCATCGCTCACCAAGGTCCTGTCAICAGATGTATACCAATGTAGAC 1967
QY 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB 1968 CAAGACCTTGTGGGTGGCGCGCTCCGCAAGGTAGCGCTCATTTGACACCTGCACTTC 2027
QY 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArg 120
DB 2028 GGTCTCTCGSACCTTTACCTGGTCAGGAGCGAGGTGATTCATTCCTGCGCGCGG 2087
QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB 2088 GGTGATAGCAGGGCAGCGCTGTGTCCGCCGCCATTCTCTACTTGAAGGCTCCTCG 2147
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaValCys 160
DB 2148 GGGGGTCCGCTGTGTGTCCCGCGGGGACCGCGTGGCATATTTAGGGCCGGGTGTC 2207
QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValLeuSerLeuGlnThrMet 180
DB 2208 ACCGTGGAGTGGTAGGGCGGTGGACTTTATCTGCTGTAGAACTATAGACACCATG 2267
QY 181 ArgSer 182
DB 2268 AGCTCC 2273

RESULT 8
AAN90336
ID AAN90336 standard: DNA: 7310 BP.
XX AC AAN90336;
XX AC
XX 25-MAR-2003 (updated)
DT 19-JUL-2001 (updated)
DT 01-NOV-1989 (first entry)
XX XX
DE Composite hepatitis C virus (HCV) cDNA.
KW Hepatitis C virus: cDNA: clone 15e; clone k9-1; probe: vaccine: ds.
XX XX
OS Pan troglodytes.
XX XX
PN GB2212511-A.
XX XX
PD 26-JUL-1989.
XX XX
PF 18-NOV-1988; 88GB-0027024.
XX XX

```

18-NOV-1987; 87US-0127114.
30-DEC-1987; 87US-0139886.
26-FEB-1988; 88US-0161072.
26-OCT-1988; 88US-0263584.
XX
PA (CHIR) CHIRON CORPORATION.
XX
PZ Houghton M, Choo QL, Kuo G;
XX
XX WPI; 1989-215054/30.
DR P-PSDB; AAP90288.
XX
PI Hepatitis C virus gene - used for prodn. of polynucleotide probes,
PT polypeptide(s) and antibodies for diagnosis, prevention and treatment
PT of infection.
XX
PS Disclosure: fig 47; 235pp; English.
XX
CC The sequence shows a composite hepatitis C virus (HCV) cDNA, derived by
CC aligning clones k9-1 through 15e in 5'-3' direction. The cDNA
CC encodes antigens which react with antibodies in patients with non-A
CC non-B hepatitis (NANBH). The cDNA can be used to design probes, or to
CC synthesize polypeptides, which are used to diagnose HCV-induced NANBH,
CC to raise antibodies for immunoassay or treatment, or to produce
CC vaccines. See also AAP90288, and AAN90303-35.
CC (N.B. This record was resubmitted to correct errors in the sequence.)
CC (Updated on 25-MAR-2003 to correct PR field.)
XX
SQ Sequence 7310 BP; 1495 A; 2218 C; 2058 G; 1539 T; 0 other;

Alignment Scores:
Pred. No.: 1,32e-77 Length: 7310
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservatives: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 10 Gaps: 0

US-09-965-594-1 (1-182) x AAN90336 (1-7310)

QY 1 MetAlaProIleThrAlaTyrAlaGlnInrArqGlyLeuLeuGlyCysLeuIleThr 20
DB 1728 CTGGCCGCCATCAGCGGTACGCCACAGACAGAGCGGCTCTAGGTGCATATCAC 1787
QY 21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyValGlnIleValSerThrAla 40
DB 1788 AGCCTAACTGGCGGACAAAACCAAGTGGAGGGTCAGTTCAGATTTGTGCAACTGCT 1847
QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAla 60
DB 1848 GCCCAAACTTCTGGCAACGTGCATCAATGGGG:GTGGTGGACTGTCTACCAACGGGGG 1907
QY 61 GlyThrArqThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
DB 1908 GGAACGAGGACCATCGCTCACCAAGGTCCTGTCAICAGATGTATACCAATGTAGAC 1967
QY 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB 1968 CAAGACCTTGTGGGTGGCGCGCTCCGCAAGGTAGCGCTCATTTGACACCTGCACTTC 2027
QY 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArg 120
DB 2028 GGTCTCTCGSACCTTTACCTGGTCAGGAGCGAGGTGATTCATTCCTGCGCGCGG 2087
QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB 2088 GGTGATAGCAGGGCAGCGCTGTGTCCGCCGCCATTCTCTACTTGAAGGCTCCTCG 2147
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaValCys 160
DB 2148 GGGGGTCCGCTGTGTGTCCCGCGGGGACCGCGTGGCATATTTAGGGCCGGGTGTC 2207
QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValLeuSerLeuGlnThrMet 180
DB 2208 ACCGTGGAGTGGTAGGGCGGTGGACTTTATCTGCTGTAGAACTATAGACACCATG 2267
QY 181 ArgSer 182
DB 2268 AGCTCC 2273

RESULT 8
AAN90336
ID AAN90336 standard: DNA: 7310 BP.
XX AC AAN90336;
XX AC
XX 25-MAR-2003 (updated)
DT 19-JUL-2001 (updated)
DT 01-NOV-1989 (first entry)
XX XX
DE Composite hepatitis C virus (HCV) cDNA.
KW Hepatitis C virus: cDNA: clone 15e; clone k9-1; probe: vaccine: ds.
XX XX
OS Pan troglodytes.
XX XX
PN GB2212511-A.
XX XX
PD 26-JUL-1989.
XX XX
PF 18-NOV-1988; 88GB-0027024.
XX XX

XX Example: Fig 16; 75pp; English.

XX The specification describes a pharmaceutical composition which comprises a hepatitis C virus (HCV) antisense polynucleotide. The HCV is characterized by a positive stranded RNA genome which has 40% homology at the polypeptide level to a HCV polypeptide. The antisense polynucleotide binds to cellular polynucleotides which enhance and/or are required for viral infectivity, replicative ability or chronicity. The antisense polynucleotides may also be designed to bind with high specificity, to be of increased stability, to be stable and to have low toxicity. The composition also comprises an agent which causes viral RNA to be inactive. The composition is used for preventing HCV replication in a system. The present sequence represents a novel HCV cDNA sequence, which is used in the course of the invention.

XX Sequence 8316 BP; 1671 A; 2529 C; 2345 G; 1771 T; 0 other:

Alignment Scores:

Pred. No.:	1:55e-77	Length:	8316
Score:	943.00	Matches:	179
Percent Similarity:	100.00%	Conservative:	3
Best Local Similarity:	98.35%	Mismatches:	0
Query Match:	98.95%	Indels:	0
DB:	21	Gaps:	0

US-09-965-594-1 (1-182) x AA75296 (1-8316)

QY : MetAlaPcTtLeThrAlaTyrAlaGlnGlnThrArgGlyGlyGlnGlyCysHisLeuPhe 20

DB 2734 CTGGGGCGGATCACCGGTACCCCGCAGACAAAGCGCTCTAGCGTGCATATACCG 2733

QY 21 SerLeuThrGlyArgAspLysAsnGlnValGlnGlyGlnValGlnLeuValSerThrAla 40

DB 2794 AGCCTAACTGGCGGGACAAAACCAAGTGGAGGTCAGGTCACAGATTGCTCAACTGCT 2853

QY 41 AlaGlnThrPheLeuAlaThrCysLeuAsnGlyValCysThrPheValThrHisLeuAla 60

DB 2854 GCCCAAACTTCCTGGCAAGTGCATCAATAGGTCTCTGGACGTGCTACCAACGCGGCC 2913

QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80

DB 2914 CGAAGAGGACCATCGCGCACCAAGGCTCTGTGATCCAGATGTATACCAAGTACAG 2973

QY 81 LysAspLeuValGlyTyrProAlaProGlnLysSerArgSerLeuThrProCysThrCys 100

DB 2974 CAAGACCTTGCGGTGGCTCGCTCGCAAGGTAGCCGTCATGACACCCGCACTTGC 3033

QY 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArgArg 120

DB 3034 GGTCTCTGGACCTTATCTGTCTACGAGGACGCGGATGTCATTCCTGGTGGCGGGG 3093

QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140

DB 3094 GTGTATACAGGGGAGCGCTGTGTGCGCGCGCGCGCATTTCCACTTGAAGGCTCCIC 3153

QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyTyrPheArgAlaAlaValCys 160

DB 3154 GGGGTGCGCTGTGTGCGCGCGCGCGCGCGCTGGGCATATTAGGCGCGCGGTGCG 3213

QY 161 ThrArgGlyValAlaLysAlaValAspPheCysLeuProValGluSerLeuLeuThrMet 180

DB 3214 ACCGTGGAGTGCTAAGCGGTGGACTTTATCCCTGGGAGCAAGCTAGACACCACTG 3273

QY 181 ArgSer 182

DB 3274 AGGTCC 3279

RESULT 11

AAZ07656

ID AAZ07656 standard; DNA; 9133 BP.

XX

AAZ07656;

20-MAR-2003 (updated)

08-NOV-1999 (first entry)

Nucleotide sequence of HCV-1 ORF.

Hepatitis C virus; HCV; J1; J7; HCV-1; non-A, non-B HCV; NANBH;

HCV infection; vaccine; ds.

Hepatitis C virus.

Key Location/Qualifiers

CDS 268..9132

/*tag= b

/transl_except= (pos:1588..1589; aa:Leu)

/note= "this codon has an apparent 1 nucleotide deletion, which alters the reading frame"

/transl_except= (pos:1647..1650; aa:Pro)

/note= "this codon has an apparent 1 nucleotide insertion, which alters the reading frame; this insertion is not indicated in the sequence present in the formal sequence listing of the specification"

EP939128-A2.

01-SEP-1999.

17-SEP-1990; 99EP-0101746.

15-SEP-1989; 89CS-0408045.

21-DEC-1989; 89CS-0456142.

17-SEP-1990; 90EP-C3-0149.

(CHIR) CHIRON CORP.

(CYAA/) OYA A.

Cha T, Han J, Houghton M, Irvine BD, Kolberg JA;

Miyamura T, Saito I, Weiner AJ;

WPI; 1599-460843/41.

P-PSDB; AAY:4375.

New Hepatitis C Virus isolates, useful for diagnosis of hepatitis infections and development of vaccines

Disclosure; Fig 12; 132pp; English.

The invention provides two new isolates of hepatitis C virus (HCV). J1 and J7. These two isolates comprise nucleotide and amino acid sequences that are distinct from the HCV isolate HCV-1. The nucleotide sequences may be used to detect non-A, non-B HCV (NANBH) polynucleotides by hybridisation for diagnosis of NANBH infections. They may also be used to screen blood donors, donated blood and blood products for this infection. The isolates may also be used to isolate other naturally occurring variants of the virus. The polypeptides may be used as a vaccine for administration to patients to protect against infection with NANBH. The present sequence represents the nucleotide sequence of HCV-1 ORF. (Updated on 20-MAR-2003 to correct PF field.)

(Updated on 20-MAR-2003 to correct PR field.)

Sequence 9133 BP; 1834 A; 2772 C; 2600 G; 1927 T; 0 other:

Alignment Scores:

Pred. NO.:	1.74e-77	Length:	9133
Score:	943.00	Matches:	179
Percent Similarity:	100.00%	Conservative:	3
Best Local Similarity:	98.35%	Mismatches:	0
Query Match:	98.95%	Indels:	0
DB:	20	Gaps:	0

US-09-965-594-1 (1-182) x AAZ07656 (1-9133)

```

QY      1 MetAlaProIleThrAlaTyrAlaGlnInThrArgGlyLeuLeuGlyCysIleIleThr 20
DB      3343 CTGGCGCCATCAGCGGCTAGCCACACACAGAGGCTCTCTAGGTCATACACACC 3402
QY      21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGluValGlnIleValSerThrAla 40
DB      3403 AGCCTAACTGGCGCGGACAAAACCAAGTGCAGGTCAGGTCGACAGATGTGTCAACTGCT 3462
QY      41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValTyrHisGlyVala 60
DB      3463 GCCCAACCTTCCTGGCAAGCTGCATCAATGGGCTGAGCTGGACTGCTACACAGGGGCC 3522
QY      61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAspValAsp 80
DB      3523 GGAAGGAGGACCATCGAGTGCACCAAGGTCCTCTCATCGAGAGTATACCAATGTAGAC 3582
QY      81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB      3583 CAAGACCTTGTGGCTGGCGCGCTCCGCAAGGAGCGGCTCATGTACACAGCTGCATTGC 3642
QY      101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArg 120
DB      3643 GGCTCCCTCGACCTTACCTGGTCACAGAGCAGCCGCAATCATTCCTGGCGCGCGCG 3702
QY      121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB      3703 GGTGATAGCAGGGCAGCGCTGCTGCCCGCGCGCCATTCTCTACTTGAAGGCTCTCTCG 3762
QY      141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
DB      3763 GGGGTCGCGCTGTGTGCGCGCGCGGCGCAGCGCTGTGGGTCATATAGGGCGCGGTGTC 3822
QY      161 ThrArgGlyValAlaLysAlaValAspPheIleProValGluSerLeuGluThrThrMet 180
DB      3823 ACCCGTGGAGTGGTAAAGCGGTGGACTTATTCCTCTGTGAGAACCTAGACAGACCACTG 3882
QY      181 ArgSer 182
DB      3883 AGGTCC 3888

RESULT 12
ID      AAQ05956 standard; DNA: 9185 HP.
AC      AAQ05956;
DT      25-MAR-2003 (updated)
DT      23-JAN-1991 (first entry)
XX      Sense strand of the compiled Hepatitis C virus cDNA sequence.
XX      Hepatitis C virus (HCV); antiviral agent; ss.
XX      Hepatitis C virus.
XX      Key Location/Qualifiers
XX      CDS 320..9185
XX      FT /*tag= a
XX      FT misc_RNA 1..1667
XX      FT /*tag= b
XX      FT /*note="epitope within this region is claimed"
XX      FT 8978..9185
XX      FT /*tag= c
XX      FT /*note="encodes an epitope that is claimed"
XX      PN EP368232-A.
XX      PD 19-SEP-1990.
XX      PF 16-MAR-1990; 9CEP-0302665.
XX      PR 18-MAY-1989; 89US-0355002.

```

```

PR      17-MAR-1989; 89US-0325338.
PR      20-APR-1989; 89US-0341334.
XX      (CHIR ) CHIRON CORP.
XX      Houghton M, Choo QL, Kuo G;
XX      WPI: 1990-284418/38.
XX      P-PSDB; AAR08124.
XX      Hepatitis C virus DNA - used for producing probes,
XX      polypeptide(s), antibodies and anti-sense polynucleotide(s) for
XX      diagnosis and therapy.
XX      Disclosure: Fig 17; 83pp; English.
XX      HCV cDNA libraries were constructed using pooled serum from a
XX      chimpanzee with chronic HCV infection. A lambda gt10 library was
XX      screened with probes derived from previously isolated clones. The
XX      ORF is derived from the overlapping clones b114a, aq30a, CA205a,
XX      CA290a, CA216a, p14a, CA167b, CA156a, CA84a, CA59a, K9-1, 253, 131,
XX      121, 141, 11b, 7f, 8h, 33c, 40b, 37b, 35, 36, 81, 32, 33b, 25c,
XX      14c, 8f, 33f, 33d, 39c, 35f, 19g, 26g, 15e, b5a and 16jh. These
XX      clones extend the sequence of the HCV genome reported in EP-38216.
XX      The upstream region from nucleotides -319 to +1348 (=11667 in this
XX      file) is covered by clones b114a, 18g, aq30a, CA205a, CA290a,
XX      CA216a, p14a, CA167b, CA156a, CA84a and CA59a; nucleotides
XX      8659-8866 (=8978-9185 in this file) are covered by clones b5a and
XX      16jh.
XX      See also AAQ05955.
XX      CC (Updated on 25-MAR-2003 to correct PA field.)
XX      SQ Sequence 9185 BP; 1849 A; 2790 C; 2608 G; 1938 T; 0 other;

Alignment Scores:
Pred. No.: 1,750-77 Length: 9185
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservative: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 11 Gaps: 0

US-09-965-594-1 (1-182) x AAQ05956 (1-9185)
QY      1 MetAlaProIleThrAlaTyrAlaGlnInThrArgGlyLeuLeuGlyCysIleIleThr 20
DB      3395 CTGGCGCCATCAGCGGCTAGCCACACACAGAGGCTCTCTAGGTCATACACC 3454
QY      21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGluValGlnIleValSerThrAla 40
DB      3455 AGCCTAACTGGCGCGGACAAAACCAAGTGCAGGTCAGGTCGACAGATTGTCAACTGCT 3514
QY      41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValTyrHisGlyVala 60
DB      3515 GCCCAACCTTCCTGGCAAGCTGCATCAATGGGCTGTGTGGACTGCTACACAGGGGCC 3574
QY      61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
DB      3575 GGAACGAGGACCATCGCTACCCACAGGGTCCTGTATCCAGATGTAATACCAATGTAGAC 3634
QY      81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB      3635 CAAGACCTTGTGGCTGGCGCGCTCCGCAAGGTCAGGTCAGGTCATTCACACCTGCACCTTC 3694
QY      101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArg 120
DB      3695 GGCTCCCTCGACCTTACCTGGTCAGAGGCGAGCCGATGTCAITCCCTGGCGCGCGG 3754
QY      121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB      3755 GGTGATAGCAGGGCAGCGCTGCTGCCCGCGCGCCATTCTCTACTTGAAGGCTCTCTCG 3814
QY      141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValIlePheArgAlaAlaValCys 160

```

```

Db 3815 GGGGTCCCGCTGTCTGCGCGGGGACGCGGTGGGCATATTAGGCGCGGGTGTGC 3874
Qy 161 ThrArgGlyValAlaLysAlaValAspPheIleProValGluSerLeuGluThrMet 180
Db 3875 ACCGTGAGTGGCTAAGGCGGTGGACTTATCCGTGTGGAGAACCTAGACACACCATG 3934
Qy 181 ArgSer 182
Db 3935 AGGTCC 3940

RESULT 13
AAQ10566
ID AAQ10566 standard; DNA; 9185 BP.
XX
AC AAQ10566:
XX
XX 25-MAR-2003 (updated)
DT 29-APR-1991 (first entry)
XX
XX Hepatitis C virus strain 1 DNA.
XX
XX Hepatitis C virus; HCV-1; non-A, non-B hepatitis; HCV antigen;
KW viral infections; ss.
XX
XX Hepatitis C virus.
OS
XX EP414475-A.
PN
XX 27-FEB-1991.
PD
XX 21-AUG-1990; 90FP-0309120.
PF
XX 25-AUG-1989; 89US-0398667.
PR
XX (CHTR ) CHIRON CORP.
PA
XX Weiner AJ, Steimer KS;
PI
XX WPI; 1991-059670/79.
DR
XX
XX Cell lines infected with hepatitis C virus - are used as source
PI of antigens for detection of HCV antibodies, for vaccines, and
PT for screening anti-viral agents
PS
XX Disclosure; fig 1; 24pp; English.
XX
XX This is a hepatitis C virus (HCV) composite cDNA sequence, deduced
CC using overlapping clones, a compo. contg. the antigenic protein
CC encoded by this sequence is useful for detecting anti-HCV anti-
CC bodies (Abs) and for screening an agent which inhibits HCV repli-
CC ation. A cell line infected with this virus can be used as a
CC source of antigens. The antigen is useful for preparing vaccines
CC for treating viral infections. See also AAQ10567.
CC (Updated on 25-MAR-2003 to correct PA field.)
XX
XX Sequence 9185 BP; 1845 A; 2790 C; 2605 G; 1938 T; 0 other;
SQ

Alignment Scores:
Pred. No.: 1.75e-77 Length: 9185
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservative: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.35% Indels: 0
DB: 12 Gaps: 0

US-09-965-594-1 (1-182) x AAQ10566 (1-9185)

Qy 1 MetAlaProIleThrAlaValAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
Db 3395 CTGGCGCCCATCACGCGGTACGCCAGACAAAGGGGCTCTCTAGGGTGCATAATCACC 3454
Qy 21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGluValGlnIleValSurThrAla 40
```

```

Db 3455 AGCCTAACTGCGCGGACAAAAACCAAGTGCAGGGTGAGGTCCAGATTGTCTCAACTGCT 3514
Qy 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpIleValIleHisGlyAla 60
Db 3515 GCCCAACCTTCCTGGCAACGTGCATCAATGGGGTGTGCTGGAGTGTCTACACGGGGCC 3574
Qy 61 GlyThrArgThrIleAlaSerProIysGlyProValIleGlnMetTyrThrAsnValAsp 80
Db 3575 GCAACGAGGACCATCGCTCACCAAGGGTCTGTCTATCCAGATGTATACCAATGTAGAC 3634
Qy 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
Db 3635 CAAGACCTTGTGGCTGGCGGCTCCGCAAGGTAGCCGCTCATTGACACACCTGCACCTGC 3694
Qy 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArg 120
Db 3695 GGCTCTCGGACCTTACCTGGTCACGAGGACCGCGATGTCATCCGCGCGCGCGG 3754
Qy 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
Db 3755 GGTATTAGCAGGGGACGCTGCTGCGCCCGGCCCATTTCTTCTACTTCAAAGGCTCCTCG 3814
Qy 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
Db 3815 GGGGTCGCTGTGTGCGCGCGGACCGCGTGGGCATATTAGGCGCGCGGTGTC 3874
Qy 151 ThrArgGlyValAlaLysAlaValAspPheIleProValGluSerLeuGluThrMet 180
Db 3875 ACCGTGAGTGGCTAAGGCGGTGGACTTATCCGTGTGGAGAACCTAGACACACCATG 3934
Qy 181 ArgSer 182
Db 3935 AGGTCC 3940

RESULT 14
AAA75297
ID AAA75297 standard; cDNA; 9185 BP.
XX
AC AAA75297:
XX
XX 15-JAN-2001 (first entry)
DT
XX Sense strand of HCV encoding a polyprotein.
DE
XX Hepatitis C virus; HCV; antisense polynucleotide; polyprotein;
KW viral infectivity; viral replication; ds.
XX
XX Hepatitis C virus.
OS
XX Key Location/Qualifiers
XX CDS 320..9184
XX /*tag= a
XX /*note= "partial sequence; no termination codon given"
XX
XX EP3034785-A2.
XX
XX 13-SEP-2000.
PD
XX
XX 16-MAR-1990; 2000EP-0109602.
PE
XX
XX 17-MAR-1989; 89US-0325338.
PR
XX 20-APR-1989; 89US-0341334.
PR
XX 18-MAY-1989; 89US-0355002.
PR
XX 16-MAR-1990; 90EP-0302866.
XX
XX (CHTR ) CHIRON CORP.
XX
XX Houghton M, Choo Q, Kuo G;
PI
XX WPI; 2000-566891/53.
DR
XX P-PSDB; AAB18541.
```


PR 10-AUG-1990; 90US-0566203.
 XX (CHTR) CHIRON CORP.
 PA Houghton M, Cheo QL, Kuo G, Weiner AJ, Gredea MS, Irvine BD;
 PI Kolberg JA;
 XX WPI: 1992-080394/10.
 DR P-PSDB; AAR21519.
 XX
 PT Reagents for isolating, amplifying and detecting HCV
 PT polynucleotide(s) - used to monitor spread of blood-borne non-A,
 PT non-B hepatitis virus infection and screen blood samples for
 PT virus
 XX
 PS Disclosure; Fig 1; 67pp; English.
 XX
 CC The sequence is a composite of HCV cDNA from HCV1, a prototypic
 CC HCV. The sequence is based upon sequence information derived from
 CC a no. of HCV cDNA clones, which were isolated from a no. of HCV
 CC cDNA libraries, including the "c" library present in lambda cll1
 CC (ATCC No.40394), and from human serum. The HCV cDNA clones
 CC were isolated by methods described in WO9014436.
 CC The clones from which the sequence was derived are 5' clones 32,
 CC b114a, 18g, aq30a, CA205a, CA290a, CA216a, p114a, CA167b, CA156e,
 CC CA84a, CA59a, K9-1 (also called K9-1), 26f, 14i, 12i, 14L, 11b, 7f,
 CC 7e, 8h, 33c, 40b, 37b, 35, 36, 81, 32, 33b, 25c, 14c, 8f, 33f, 33g,
 CC 29c, 35f, 19a, 26g, 15e, b5a, 16jh, 6k and p31j3.
 CC The target regions indicated in the features are listed in claim 1,
 CC page 49. Oligomers are provided which are complementary to these
 CC target regions and used in the detection of an HCV sequence in an
 CC analyte.
 CC (Updated on 25-MAR-2003 to correct PA field.)
 CC (Updated on 25-MAR-2003 to correct P1 field.)
 XX
 SQ Sequence 9400 BP: 1385 A: 2460 C: 2671 G: 1964 T: 0 other:

Search completed: September 27, 2003, 12:36:32
 Job time : 344 secs

Alignment Scores:
 Pred. No.: 1-810-77 Length: 9400
 Score: 943.00 Matches: 179
 Percent Similarity: 100.00%
 Best Local Similarity: 100.00%
 Query Match: 98.35%
 Mismatches: 0
 Indels: 0
 Gaps: 0
 DB: 13
 US-09-965-594-1 (1-182) x AAQ2:744 (1-9436)
 QY 1 MetAlaProfilerThrAlaTyAlaAlaGlnThrArgGlyLeuLeuGlyCysIleGlnThr 20
 Db 3416 CTGGCCGCCATACGGCGTACGCCACGACAGCAAGGGGCGTCTGTAGGTGCATAATCAGC 3475
 QY 21 SerLeuThrGlyArgAspLysAsnGlnValGlnGlyGlyValGlnIleValSerThrAla 40
 Db 3476 AGCCTAACCTGGCGGGGACAAAACCAAGTGGAGGGTGGAGTGTGCAGATGTGTCAACTGCT 3535
 QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValTyHisGlyVala 60
 Db 3536 GCCCAACCTTCTGCGACAGGTGCATCATGGGTGTGTGACATGTCTACCAAGGGGCGG 3595
 QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyThrAsnValAsp 80
 Db 3596 GGAACAGGACCATCCGTCACCAAGGGTGTGTGTCATCAGATGTATACCAATGTAGAC 3655
 QY 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
 Db 3656 CAAGACCTTGTGGGTGGCGCGCTCCGAAGGTAGCGGTGATTCACACCTGCACITGC 3715
 QY 101 GlySerSerAspLeuTyLeuValThrArgHisAlaAspValIleProValArgArgArg 120
 Db 3716 GGCCTCTCGGAGCCTTACCTGTGTACAGAGGACAGCGCATGTCTATTCCTGCGCGCGG 3775
 QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyLeuLysGlySerSer 140
 Db 1401 GGTGCTCTGCGGAGCCTTACCTGTGTACAGAGGACAGCGCATGTCTATTCCTGCGCGG

GenCore version 5.1.6
Copyright: (c) 1993 - 2003 Computer Aid.

OM protein - nucleic search, using frame_plus_p2n model

Run on: September 27, 2003, 12:22:50 : Search time 3348 Seconds
(without alignments)
2223.881 Million cell updates/sec

Title: US-09-965-594-1
Perfect score: 953
Sequence: : MAP::YAAQTRGL:GCIT.....GVAKAVDFIPVSLDT:MRS 182

Scoring table: BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 2668711 seqs, 2045483396 residues

Total number of hits satisfying chosen parameters: 577422

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 99%
Listing first 45 summaries

Command line parameters:

-MODEL=frame_plus_model -DEV=xih
-Q/cgn2_1/uspto.spoc1/us09965594/runat_26092003_164343_18756/app_query.fasta_1.327
-DB=GenEmbl -FAST=fastap -SUFFIX=rge -M=MMATCH=0.1 -LCOPT=0 -LCOPEXT=0
-UNITS=bits -START=1 -END=1 -MA=RIX-blosum62 -TRANS=human40.csi -LIST=45
-DOCAT=GN=200 -THR SCORE=ect -THR MAX=99.9 -THR MIN=0 -AL=GN=15 -MODE=LOCAL
-OUTEXT=ptc -NORM=ext -HEARSIZE=500 -MINLEN=0 -MAXLEN=200000000
-USER=US09965594 -CGN_1_1_3508#runat_26092003_164343_18756 -NGPU=6 -ICPU=3
-NO_MMAP -LARGEQUERY -NEG.SCORES=0 -WAIT -DSPEX=CC=160 -LONGLOG
-DSV_TIMEOUT=120 -WARN_TIMEOUT=30 -THREAFS=1 -XGAPOP=10 -XGAPEXT=0.5 -Fgapop=6
-Fgapext=7 -Ygapop=10 -Ygapext=0.5 -Delop 6 -Delext=7

Database :

GenEmbl:

1: gb_ba:*
2: gb_hgt:*
3: gb_in:*
4: gb_om:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vi:*
15: em_ba:*
16: em_fun:*
17: em_hum:*
18: em_in:*
19: em_mu:*
20: em_om:*
21: em_or:*
22: em_ov:*
23: em_pat:*
24: em_ph:*
25: em_pl:*
26: em_ro:*
27: em_sts:*
28: em_un:*

29: em_vir:*
30: em_htg_hum:*
31: em_htg_inv:*
32: em_htg_other:*
33: em_htg_mus:*
34: em_htg_pln:*
35: em_htg_rod:*
36: em_htg_mam:*
37: em_htg_vrt:*
38: em_sy:*
39: em_htgo_hum:*
40: em_htgo_mus:*
41: em_htgo_other:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	946	99.3	2058	6	AX395309 Sequence
2	946	99.3	2058	6	AX454818 Sequence
3	945	99.2	543	14	AF369218 Hepatitis
4	945	99.2	543	14	AF369235 Hepatitis
5	943	99.0	5360	6	AR118686 Sequence
6	943	99.0	5360	6	I06434 Sequence 48
7	943	99.0	5360	6	I03328 Sequence 8
8	943	99.0	6785	6	AR118692 Sequence
9	943	99.0	6785	6	I06440 Sequence 54
10	943	99.0	6785	6	I03329 Sequence 10
11	943	99.0	7310	6	AR118696 Sequence 15
12	943	99.0	7310	6	I09331 Sequence 15
13	943	99.0	7310	14	HPCPOLYP M32084 Hepatitis C
14	943	99.0	8316	6	AR118703 Sequence
15	943	99.0	8987	6	AR118728 Sequence
16	943	99.0	9185	6	AR118722 Sequence
17	943	99.0	9185	6	AR118723 Sequence
18	943	99.0	9185	6	BD091382 HCV cult1
19	943	99.0	9185	6	I04254 Sequence 1
20	943	99.0	9379	6	AR166930 Sequence
21	943	99.0	9379	6	AR301300 Sequence
22	943	99.0	9401	6	AR176483 Sequence
23	943	99.0	9401	6	BD080334 Hepatitis
24	943	99.0	9401	6	E65593 Hepatitis C
25	943	99.0	9401	6	I71894 Sequence 9
26	943	99.0	9401	6	I81885 Sequence 9
27	943	99.0	9401	14	HPCPLYPRE M62321 Hepatitis C
28	943	99.0	9409	12	AF387805 Synthetic
29	943	99.0	9409	12	AF387808 Synthetic
30	943	99.0	9418	14	AF274632 Hepatitis
31	943	99.0	9418	14	AF387806 Synthetic
32	943	99.0	9693	12	AF387807 Synthetic
33	942	98.8	543	14	AF369222 Hepatitis
34	942	98.8	543	14	AF369232 Hepatitis
35	942	98.8	543	14	AF369240 Hepatitis
36	942	98.8	543	14	AF369245 Hepatitis
37	942	98.8	2061	6	AX441176 Sequence
38	942	98.8	2061	6	AX457113 Sequence
39	940	98.6	543	14	AF369224 Hepatitis
40	940	98.6	543	14	AF369237 Hepatitis
41	940	98.6	9424	14	AF511948 Hepatitis
42	939	98.5	543	14	AF369230 Hepatitis
43	939	98.5	9502	6	E08263 gRNA of Hep
44	939	98.5	9502	6	E08264 cDNA of Hep
45	939	98.5	9502	14	HFCHCJ1 D10749 Hepatitis C

ALIGNMENTS

RESULT 1

```

AX395309
LOCUS AX395309 2058 bp DNA linear PAT 18-MAY-2002
DEFINITION Sequence 2 from Patent WO0196875.
ACCESSION AX395309
VERSION AX395309.1 GI:21066308
KEYWORDS
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE
1 Chien,D.Y., Arcangel,P., Tandeske,L., George-Nascimento,C.,
AUTHORS Coit,D. and Medina-Solby,A.
TITLE Hcv antigen/antibody combination assay
JOURNAL Patent: WO 0196875-A 20-DEC-2002
CHIRON CORPORATION (US)
FEATURES
Location/Qualifiers
source
1..2058
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/notes="representative NS3/4a conformational antigen"
1..>2058
/notes="unnamed protein product"
/codon_start=1
/transl_table=11
/protein_id="CAD32154.1"
/db_xref="GI:21066309"
/translation="MAPITAYAOQTRGLGCIISLTGRDKNQVEGEVQIVSTAAQTF
LATCINGVCTVYHGAGRTIASPKGPVIOYINVDOLVGMFAFGSRSLPTCTGSS
SDLYLVTRHAPVIVREBGSBGLSDPLISYKGGSGGPLLCHNAGVAFRAVC
TRGVAKAVDFIPVENLETMRSPVFDNSPPVPOFQVHLHAPTSKSKSVAAA
YAAQGYKVLNPSVAATLFGAYMSKANGIDNRTGVRITIGSPIITSTYKFLA
DGGSGGAYDIICDECHSDATSLGIGTVLQAGTAGAPHLVLAATATPGSVVPH
PNIIEVALSTTGEIPFYKAIPLVIGKGRRLPCRKKKCDLAAKLALGAINAVAY
YRGLDVSVIPDGVVVAATDALMTGYTGDFSDVDCNTCVTQVDFSLDPTFTET
TLPOQAVSRTOGRGKPGIYREVAQGPSPGDFSSVLCGECYDAGCAWYELTPA
ETTVRLAYMNPGLPCVDHLEFNEGVTGLTHDAHFLSOTKSGENLPLVAYQA
TVCARACAPPSNDQMKCLIRKPTLHGPTPLLYRLGAVQNEITLTHPTVYIMTCM
SADLEVYTSWLVGGVLAALAAAYCLSTGCVVIVGRVVSCKPAIIPDREVLVREFDE
MEEC"
BASE COUNT 419 a 634 c 580 g 425 t
ORIGIN
Alignment Scores:
Pred. No.: 6,56e-69 Length: 2058
Score: 946.00 Matches: 180
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 98.90% Mismatches: 0
Query Match: 99.27% Indels: 0
DB: 6 Gaps: 0
US-09-965-594-1 (1-182) x AX395309 (1-2058)
Qy 1 MetAlaProIleThrAlaIleThrArgGlyLeuLeuGlyCysIleIleThr 20
Db 1 ATGGCGCCATACAGGCGTACCGCCAGCAGACAGAGGCGCTCTAGGTCGATATCAGC 60
Qy 21 SerLeuThrGlyArgAspIleAsnGlnValLeuGlyCysValGlnIleValSerThra 40
Db 61 AGCCTAACTGGCGCGGGAACAAACCAACTGAGGCTGAGTCCAGATGTCGTCACTGCT 120
Qy 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTyrIleThrValTyrHisGlyAla 60
Db 121 CCCCAACCTCTCTGGCAACGTGCATCAATCGGTCGCTGCGAATGCTACCAAGGCGCC 180
Qy 61 GlyThrArgThrIleAlaSerProGlyGlyProValIleGlnMetTyrThrAsnValAsp 80
Db 181 GGAACGAGGACCATCGCTCACCCAAAGGCTCTGTCTCATCCAGATGCTATACCAATGTAGC 240
Qy 81 LysAspLeuValGlyTrpProIleProGlnGlySerArgSerLeuThrProCysThrCys 100
Db 241 CAAGACCTTGTGGGTGGCGCGGCTCCGCAAGGTAGCCGATATTACACCCCTGCAC-TGC 300

```

```

Qy 101 GlySerSerAspLeuTyrIleuValThrArgHisAlaAspValIleProValArgArgArg 120
Db 301 GGCTCTCGACCTTTACCTGTGTACAGGACGACGCGATGTCAITCCCGTGGCGCGCG 360
Qy 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
Db 361 GGTGATAGCAGGCGACGCTGTGTGCGCGCGGCCCATTTCTACTTGAAGGCTCCTCG 420
Qy 141 GlyGlyProLeuLeuCysProAagGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
Db 421 GGGGCTCCCTGTGTGTGCGCGCGGCGACGCGTGGCATATTTAGGCGCGCGGTGTC 480
Qy 161 ThrArgGlyValAlaLysAlaValAspPheIleProValIleLeuSerLeuGlnThrMet 180
Db 481 ACCCGTGGAGTGGCTAAGCGGTGGACTTTATCCCTGTGGAGAACCTAGACAGCAACATG 540
Qy 181 ArgSer 182
Db 541 AGGTCC 546
RESULT 2
AX454818
LOCUS AX454818 2058 bp DNA linear PAT 06-JUL-2002
DEFINITION Sequence 1 from Patent WO0196870.
ACCESSION AX454818
VERSION AX454818.1 GI:21714047
KEYWORDS
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE
1 Chien,D.Y., Arcangel,P., Tandeske,L., George-Nascimento,C.,
AUTHORS Coit,D. and Medina-Solby,A.
TITLE Immunoassays for anti-hcv antibodies
JOURNAL Patent: WO 0196870-A 20-DEC-2002;
CHIRON CORPORATION (US)
FEATURES
Location/Qualifiers
source
1..2058
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/notes="representative NS3/4a conformational antigen"
1..>2058
/notes="unnamed protein product"
/codon_start=1
/transl_table=11
/protein_id="CAD38232.1"
/db_xref="GI:21714048"
/translation="MAPITAYAOQTRGLGCIISLTGRDKNQVEGEVQIVSTAAQTF
LATCINGVCTVYHGAGRTIASPKGPVIOYINVDOLVGMFAFGSRSLPTCTGSS
SDLYLVTRHAPVIVRRGDSRGSLLSPISYKGGSGGPLLCHNAGVAFRAVC
TRGVAKAVDFIPVENLETMRSPVFDNSPPVPOFQVHLHAPTSKSKSVAAA
YAAQGYKVLNPSVAATLFGAYMSKANGIDNRTGVRITIGSPIITSTYKFLA
DGGSGGAYDIICDECHSDATSLGIGTVLQAGTAGAPHLVLAATATPGSVVPH
PNIIEVALSTTGEIPFYKAIPLVIGKGRHLIFCHSKKCDLAAKLALGAINAVAY
YRGLDVSVIPDGVVVAATDALMTGYTGDFSDVDCNTCVTQVDFSLDPTFTETI
TLPOQAVSRTOGRGKPGIYREVAQGPSPGDFSSVLCGECYDAGCAWYELTPA
ETTVRLAYMNPGLPCVDHLEFNEGVTGLTHDAHFLSOTKSGENLPLVAYQA
TVCARACAPPSNDQMKCLIRKPTLHGPTPLLYRLGAVQNEITLTHPTVYIMTCM
SADLEVYTSWLVGGVLAALAAAYCLSTGCVVIVGRVVSCKPAIIPDREVLVREFDE
MEEC"
BASE COUNT 419 a 633 c 581 g 425 t
ORIGIN
Alignment Scores:
Pred. No.: 6,56e-68 Length: 2058
Score: 946.00 Matches: 180
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 98.90% Mismatches: 0
Query Match: 99.27% Indels: 0
DB: 6 Gaps: 0
US-09-965-594-1 (1-182) x AX454818 (1-2058)

```


Db 1470 AGGTCC 1475

RESULT 8
AR118692

LOCUS AR118692 6785 bp DNA linear PAT 16-MAY-2001
DEFINITION Sequence 65 from patent US 6150087.
ACCESSION AR118692
VERSION AR118692.1 GI:14100602
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 6785)
AUTHORS Chien,D.Y.
TITLE MANV diagnostics and vaccines
JOURNAL Patent: US 6150087-A 65 21-MAY-2000;
FEATURES Location/Qualifiers
source 1..6785 /organism="unknown"
BASE COUNT 1392 a 2050 c 1914 g 1429 t
ORIGIN

Alignment Scores:
Pred. No.: 4.15e-67 Length: 6785
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservative: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
Gaps: 0
DB:

US-09-965-594-1 (1-192) x AR118692 (1-6785)

QY 1 MetAlaProIleThrAlaTyAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
DB 1203 CTGGCGCCCATCACGGCTACGCCAGCAGACAGAGGGCCCTCTAGGCTGCATAATCAC 1262
QY 21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGluValGlnIleValSerThrAla 40
DB 1263 AGCCTAACTGGCGGGGACAAAACCAAGTGGAGGTGAGGTCCAGATTGTCTCAACTGCT 1322
QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTyrThrValTyrHisGlyAla 60
DB 1323 GCCCAAACTTCCTGGCAACGTGCATCAATGGGTGTCTGGACTGTCTACACGGGGCC 1382
QY 61 GlyThrArgIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 90
DB 1383 GGAACGAGGACCATCGCTCACCCAGAGTCTCTGCATCCAGATGTAACCAATGTACAC 1442
QY 81 LysAspLeuValGlyTyrProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB 1443 CAAGACCTTGTGGCTGGCGGCTCCGCAAGGTAGCGGCTCATTGACACCTGCACCTGC 1502
QY 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
DB 1503 GGCTCTCGGACCTTACCTGGTCCAGGACCGCGCATGTCATTCCTGCTGGCGGGCGG 1562
QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB 1563 GGTGATAGAGGGGACGCTGCTGCCCGGGCCCATTTCTTCTACTTGAAGGCTCCTCG 1622
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
DB 1623 GGGGCTCCGCTGTGTGGCGGGGACGCGGTGGGCAATATTAGGGCGGGGTGTGC 1682
QY 161 ThrArgGlyValAlaAlaLysAlaValAspPheIleProValGluSerLeuGluThrThrMet 180
DB 1683 ACCCGTGGAGTGGCTAAGCGGGTGGACTTTATCCCTGTGGAGAACCTAGACACCAACATG 1742
QY 181 ArgSer 182
DB 1743 AGGTCC 1748

RESULT 10
109329

LOCUS 109329 6785 bp DNA linear PAT 02-DEC-1994
DEFINITION Sequence 10 from Patent WO 8904669.

106440

LOCUS 106440 6785 bp DNA linear PAT 02-DEC-1994
DEFINITION Sequence 54 from Patent EP 0318216.
ACCESSION 106440
VERSION 106440.1 GI:590312
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 6785)
AUTHORS Houghton,M., Choo,Q.-L. and Kuo,G.
TITLE Nanbv diagnostics and vaccines
JOURNAL Patent: EP 0318216-A1 54 31-MAY-1989;
FEATURES Location/Qualifiers
source 1..6785 /organism="unknown"
BASE COUNT 1392 a 2050 c 1914 g 1429 t
ORIGIN

Alignment Scores:
Pred. No.: 4.15e-67 Length: 6785
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservative: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
Gaps: 0
DB:

US-09-965-594-1 (1-182) x 106440 (1-6785)

QY 1 MetAlaProIleThrAlaTyAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
DB 1203 CTGGCGCCCATCACGGGTACGCCAGCAGACAGAGGGCCCTCTAGGCTGCATAATCAC 1262
QY 21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGluValGlnIleValSerThrAla 40
DB 1263 AGCCTAACTGGCGGGGACAAAACCAAGTGGAGGTGAGGTCCAGATTGTCTCAACTGCT 1322
QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTyrThrValTyrHisGlyAla 60
DB 1323 GCCCAAACTTCCTGGCAACGTGCATCAATGGGTGTCTGGACTGTCTACACGGGGCC 1382
QY 61 GlyThrArgIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
DB 1383 GGAACGAGGACCATCGCTCACCCAGAGTCTCTGCATCCAGATGTAACCAATGTACAC 1442
QY 81 LysAspLeuValGlyTyrProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB 1443 CAAGACCTTGTGGCTGGCGGCTCCGCAAGGTAGCGGCTCATTGACACCTGCACCTGC 1502
QY 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
DB 1503 GGCTCTCGGACCTTACCTGGTCCAGGACCGCGCATGTCATTCCTGCTGGCGGGCGG 1562
QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB 1563 GGTGATAGAGGGGACGCTGCTGCCCGGGCCCATTTCTTCTACTTGAAGGCTCCTCG 1622
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
DB 1623 GGGGCTCCGCTGTGTGGCGGGGACGCGGTGGGCAATATTAGGGCGGGGTGTGC 1682
QY 161 ThrArgGlyValAlaAlaLysAlaValAspPheIleProValGluSerLeuGluThrThrMet 180
DB 1683 ACCCGTGGAGTGGCTAAGCGGGTGGACTTTATCCCTGTGGAGAACCTAGACACCAACATG 1742
QY 181 ArgSer 182
DB 1743 AGGTCC 1748

RESULT 10
109329

LOCUS 109329 6785 bp DNA linear PAT 02-DEC-1994
DEFINITION Sequence 10 from Patent WO 8904669.

```

ACCESSION      I09329
VERSION        I09329.1  GI:587964
KEYWORDS      (unknown)
SOURCE        Unknown.
ORGANISM      Unclassified.
REFERENCE     1 (bases 1 to 6785)
AUTHORS      Broughton, M., Choo, Q.-K., and Kuo, G.
JOURNAL      Patent: WO 8904689-A 10.01-JUN-1989;
FEATURES     Location/Qualifiers
             source          1..6785
             BASE COUNT     1392 a 2050 c 1914 g 1423 t
             ORIGIN
Alignment Scores:
Pred. No.:      4,156-67      Length:      6785
Score:          943.00      Matches:      179
Percent Similarity: 100.00%      Conservative: 3
Best Local Similarity: 98.35%      Mismatches: 0
Query Match:    98.95%      Indels:      0
DB:             6      Gaps:      0
US-09-965-594-1 (1-182) x I09329 (1-6785)
QY      1 MetAlaProIleThrAlaTyrAlaGlnGlnThrArgGlyLeuLeuGlyCysLeileIleThr 20
DB      1203 CTGGGGCCCATCACGGCGTACGCCAGCAGACAGAGGGGCTCTCTAGGGTGCATATACACC 1262
QY      21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGlnValGlnIleValSerThrAla 40
DB      1263 AGCCTAATCGCGGGACAAAACCAACCAAGTGGAGGTGAGTCCAGATTGTGTCAACTGCT 1322
QY      41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAla 60
DB      1323 GCCCAAACTCTCTGCAACGTCATCAATATGAGGCTGTGATGACTGTCTACCAAGGGGCC 1382
QY      61 GlyThrArgThrIleAlaSerProLysGlyProValIleLeuMetTyrThrAsnValAsp 80
DB      1393 GGAACGAGACCATCCGCTACACCAAGGTCCTGTCATCCAGATGATACCAATGTAGAC 1442
QY      81 LysAspLeuValClyTrpProAlaProGlnGlySerArgSerSerSerThrProCysThrCys 100
DB      1443 CAAGACCTTGTGGGCTGGCGGCTCGGCAAGGATGAGCGGTCTATGACACCTGGCACTTGC 1502
QY      121 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
DB      1503 GGCTGTCTGGAGCTTTACTTGGTCAAGAGGACAGCGGATGTCTATTCCTCGCCCGGGG 1562
QY      121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB      1563 GGTCAATAGCAGGGGACAGCTGTGTGTCGGCGCGGAGATTCCTACTGTGAAGACTCTGTC 1622
QY      141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
DB      1623 GGGGGTCCGCTGTGTGGCGCGGCGGACAGCGCGCTGGCAATATTTAGGCGCGGTGTGC 1682
QY      161 ThrArgGlyValAlaLysAlaValAspPheIleProValGluSerLeuGluThrThrMet 180
DB      1683 ACCCGTGGAGTGGCTAAGCGGTGGGACTTATCTCCGTGTGAGAACCTAGAGATAACCAATG 1742
QY      181 ArgSer 182
DB      1743 AGGTCC 1748
RESULT 11
LOCUS      AR118696              7310 bp      DNA              linear      PAT 16-MAY-2001
DEFINITION Sequence 74 from patent US 6150087.
ACCESSION AR118696
VERSION   AR118696.1  GI:14106606
KEYWORDS  (unknown)
SOURCE    Unknown.
REFERENCE 1 (bases 1 to 7310)
LOCUS      T09331              7310 bp      DNA              linear      PAT 02-DEC-1994
DEFINITION Sequence 15 from Patent WO 8904669.
ACCESSION T09331
VERSION   T09331.1  GI:587966
KEYWORDS  (unknown)
SOURCE    Unknown.
ORGANISM  Unclassified.
REFERENCE 1 (bases 1 to 7310)
ORGANISM  Unknown.
REFERENCE 1 (bases 1 to 7310)

```



```

1728 CTGGGCGCCATACCGGCTACGCGCAGCAAGAGGCGCTCTAGGGGCGCATATACCG 1782
QY
21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGlnValGlnIleValSerThrAla 60
1788 AGCCCAACCTGGCGGCGGCAAAACCAAGTGGAGGCTGAGGTCACAGATGTGTCAACTCT 1847
QY
41 AlaGlnThrProLeuAlaThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAla 60
1848 GCCCAAACTTCTGGCAACGTCATCAATGGGCTGTCTGACTGTCTACGACACGGGCG 1907
DB
61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
1908 GGAACAGCAGCACCATCGCTACCCCAAGGCTCTGTCAATCCAGATGATACCAATATGAC 1967
QY
81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
1968 CAAGACCTTGTGGGCTGGCGCGCTCGGCAAGGTCTGTCAATCCAGATGATACCAATATGAC 2027
DB
101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
2028 GGCCTCTCGGACCTTACCTGTGTACAGGCGCAGCGCGATGTCTATCCCTGGCGGCGG 2087
DB
121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
2088 GGTGATAGCAGGGCAGCGTGTGTGGCGCGCGCCCATTTCTACTTGAAGAGCTCTCTCG 2147
QY
141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
2148 GGGGTCCTGCTGTGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 2207
DB
161 ThrArgGlyValAlaAlaLysAlaValAspPheIleProValGluSerLeuGluThrThrMet 180
2208 ACCGTGTGAGTGGCTAAGCGGTGGACTTTATCCCTGTGGAGACCTAGAGACACCAATG 2267
QY
181 ArgSer 182
2268 AGGTCC 2273
DB
RESULT 14
AR118703
LOCUS AR118703 8316 bp DNA linear PAT 16-MAY-2001
DEFINITION Sequence 88 from patent US 6150087.
ACCESSION AR118703
VERSION AR118703.1 GI:14100613
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 8316)
AUTHORS Chien,D.Y.
TITLE NANBV diagnostics and vaccines
JOURNAL Patent: US 6150087-A 88 21-NOV-2000;
FEATURES
Source location/Qualifiers
Source 1..8316 /organism="unknown"
BASE COUNT 1671 a 2529 c 2645 g 2771 t
ORIGIN
Alignment Scores:
Pred. No.: 5,17e-67 Length: 8316
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservative: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 6 Gaps: 0
US-09-965-594-1 (1-182) x AR118703 (1-8316)
QY
1 MetAlaProIleThrAlaTyrAlaGlnGlnThrArgGlyLeuLeuGlyCysIleThr 20
2734 CTGGCGCCCATCAGCGGTAGCGCCAGCAGACAAAGGGCGCTCTAGGGTGCATATCAC 2793
DB

```

```

21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGlnValGlnIleValSerThrAla 40
2794 AGCCCAACCTGGCGGCGGCAAAACCAAGTGGAGGCTGAGGTCACAGATGTGTCAACTCT 2853
QY
41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAla 60
2854 GCCCAAACTTCTGGCAACGTCATCAATGGGCTGTCTGACTGTCTACGACACGGGCG 2913
DB
61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
2914 GGAACAGCAGCACCATCGCTACCCCAAGGCTCTGTCAATCCAGATGATACCAATATGAC 2973
DB
81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
2974 CAAGACCTTGTGGGCTGGCGCGCTCGGCAAGGTATGCCGCTCATTCGACACCTGCACTTC 3033
DB
101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
3034 GGCCTCTCGGACCTTACCTGTGTACAGGCGCAGCGCGATGTCTATCCCTGGCGGCGG 3093
DB
121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
3094 GGTGATAGCAGGGCAGCGTGTGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 3153
DB
141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
3154 GGGGTCCTGCTGTGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 3213
QY
161 ThrArgGlyValAlaAlaLysAlaValAspPheIleProValGluSerLeuGluThrThrMet 180
3214 ACCGTGTGAGTGGCTAAGCGGTGGACTTTATCCCTGTGGAGACCTAGAGACACCAATG 3273
DB
181 ArgSer 182
3274 AGGTCC 3279
DB
RESULT 15
AR118728
LOCUS AR118728 8987 bp DNA linear PAT 16-MAY-2001
DEFINITION Sequence 137 from patent US 6150087.
ACCESSION AR118728
VERSION AR118728.1 GI:14100638
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 8987)
AUTHORS Chien,D.Y.
TITLE NANBV diagnostics and vaccines
JOURNAL Patent: US 6150087-A 137 21-NOV-2000;
FEATURES
Source location/Qualifiers
Source 1..8987 /organism="unknown"
BASE COUNT 1807 a 2735 c 2547 g 1898 t
ORIGIN
Alignment Scores:
Pred. No.: 5,62e-67 Length: 8987
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservative: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 6 Gaps: 0
US-09-965-594-1 (1-182) x AR118728 (1-8987)
QY
1 MetAlaProIleThrAlaTyrAlaGlnGlnThrArgGlyLeuLeuGlyCysIleThr 20
3076 CTGGCGCCCATCAGCGGTAGCGCCAGCAGACAAAGGGCGCTCTAGGGTGCATATCAC 3135
DB
21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGlnValGlnIleValSerThrAla 40
3136 AGCTTAACCTGGCGGCGGCAAAACCAAGTGGAGGCTGAGGTCAGATTTGTCAACTGCT 3195
DB

```

```
QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAla 60
Db 3196 GCCCAACCTTCCCTGGCAACATGCATCAATGGGTGGCTGGACTGTCTACCCAGGGGGCC 3255
QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
Db 3256 GGAACGAGGACCATCGGTCAACCAAGGTCCTGTCTCATCCAGATGATACCAATGTAGAC 3315
QY 81 LysAspLeuValGlyTyrProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
Db 3316 CAAGACCTTGTGGGCTGGCCGCTCCGCCAAGTAGCCGCTATTGACACCTGCACCTGG 3375
QY 101 GlySerSerAspLeuValThrArgHisAlaAspValIleProValArgArgArg 120
Db 3376 GCTCTCTCGGACCTTACCTGCTCAGCAGGACGCGCATGTCTACTTGAAAGGCTCTGG 3435
QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrTyrLeuLysGlySerSer 140
Db 3436 GGTGATACGAGGACGCTACTGTCTGCTCCGCGGCTCTCTACTTGAAAGGCTCTGG 3495
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheAsnAlaAlaValCys 160
Db 3496 GGGGSHYGGCTTGTGTCGCGCGGACGCGTGTGSGCATATTAGGSCATAGSTGTGG 3555
QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValIleIleSerLeuSerThrThrMet 180
Db 3556 ACCGTGAGTGGCTAAGCGGTGSACTTTATGCTGTGAGAACCTAGAGACACCATG 3615
QY 181 ArgSer 182
Db 3616 AGGTCC 3621
```

Search completed: September 27, 2003, 13:32:26
Job time : 3361 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 Computer Ltd.

OM protein - protein search, using sw model

Run on: September 27, 2003, 11:35:53 : Search time 107 Seconds
(without alignments)
336.931 Million cell updates/sec

Title: US-09-965-594-1

Perfect score: 953

Sequence: 1 MAPITAYAQTRGLGCIIT.....GVAXAVDFIPVESLEIMKPS 182

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 6

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 99%

Listing first 45 summaries

Database :

SPREMBL_23.1:

- 1: sp_archaea:
- 2: sp_bacteria:
- 3: sp_fungi:
- 4: sp_human:
- 5: sp_invertebrate:
- 6: sp_mammal:
- 7: sp_mbc:
- 8: sp_oragaele:
- 9: sp_phage:
- 10: sp_plant:
- 11: sp_rodent:
- 12: sp_virus:
- 13: sp_vertebrate:
- 14: sp_unclassified:
- 15: sp_virus:
- 16: sp_bacteriap:
- 17: sp_archaeap:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	945	99.2	181	12 Q91RS8	Q91RS8 hepatitis C
2	945	99.2	181	12 Q91RT5	Q91RT5 hepatitis C
3	943	99.0	2436	12 Q81756	Q81756 hepatitis C
4	943	99.0	3011	12 Q91FES	Q91FES hepatitis C
5	942	98.8	181	12 Q91BR3	Q91BR3 hepatitis C
6	942	98.8	181	12 Q91RS1	Q91RS1 hepatitis C
7	942	98.8	181	12 Q91RQ8	Q91RQ8 hepatitis C
8	942	98.8	181	12 Q91RT1	Q91RT1 hepatitis C
9	940	98.6	181	12 Q91R36	Q91R36 hepatitis C
10	940	98.6	181	12 Q91RS9	Q91RS9 hepatitis C
11	939	98.5	181	12 Q91RS3	Q91RS3 hepatitis C
12	939	98.5	3011	12 Q03463	Q03463 hepatitis C
13	938	98.4	181	12 Q91RT4	Q91RT4 hepatitis C
14	938	98.4	181	12 Q91RS8	Q91RS8 hepatitis C
15	938	98.4	181	12 Q91RT3	Q91RT3 hepatitis C
16	938	98.4	181	12 Q91RS5	Q91RS5 hepatitis C

17	938	98.4	181	12 Q91RS7	Q91RS7 hepatitis C
18	938	98.4	181	12 Q91RT0	Q91RT0 hepatitis C
19	937	98.3	3011	12 Q36579	Q36579 hepatitis C
20	936	98.2	181	12 Q91RS5	Q91RS5 hepatitis C
21	936	98.2	181	12 Q91RS4	Q91RS4 hepatitis C
22	935	98.1	181	12 Q91RT6	Q91RT6 hepatitis C
23	934	98.0	181	12 Q91RT9	Q91RT9 hepatitis C
24	933	97.9	181	12 Q91RS4	Q91RS4 hepatitis C
25	933	97.9	181	12 Q91RS9	Q91RS9 hepatitis C
26	933	97.9	181	12 Q91RS0	Q91RS0 hepatitis C
27	933	97.9	3011	12 Q91RS9	Q91RS9 hepatitis C
28	932	97.8	181	12 Q91RS2	Q91RS2 hepatitis C
29	931	97.7	181	12 Q91RR7	Q91RR7 hepatitis C
30	931	97.7	3011	12 Q91RT6	Q91RT6 hepatitis C
31	930	97.6	181	12 Q91RT2	Q91RT2 hepatitis C
32	930	97.6	181	12 Q91RS1	Q91RS1 hepatitis C
33	930	97.6	181	12 Q91RS2	Q91RS2 hepatitis C
34	930	97.6	181	12 Q91RS2	Q91RS2 hepatitis C
35	930	97.6	3011	12 Q36608	Q36608 hepatitis C
36	930	97.6	3015	12 Q9PWK5	Q9PWK5 hepatitis C
37	930	97.6	3015	12 Q9PWJ9	Q9PWJ9 hepatitis C
38	928	97.4	181	12 Q91RS6	Q91RS6 hepatitis C
39	927	97.3	181	12 Q91RT7	Q91RT7 hepatitis C
40	925	97.1	181	12 Q91RS0	Q91RS0 hepatitis C
41	925	97.1	181	12 Q91RT8	Q91RT8 hepatitis C
42	924	97.0	3011	12 Q36609	Q36609 hepatitis C
43	919	96.4	3011	12 Q36610	Q36610 hepatitis C
44	909	95.4	3011	12 Q81754	Q81754 hepatitis C
45	903	94.8	3010	12 Q9J3G9	Q9J3G9 hepatitis C

ALIGNMENTS

RESULT 1

Q91RS8
ID Q91RS8 PRELIMINARY: PRT: 181 AA.
AC Q91RS8:
BT 01-DEC-2001 (TRENBLrel, 19, Created)
BT 01-DEC-2001 (TRENBLrel, 19, Last sequence update)
BT 01-MAR-2003 (TRENBLrel, 23, Last annotation update)
DE NS3 protease (Fragment).
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID:11103;
RN 111
RF SEQUENCE FROM N.A.
RC STRAIN-Pt.1Y;
KA Holland-Staley C.A., Kovari L.C., Golenberg E., Mayers D.L.;
RT "Genetic diversity and response to IFN of the NS3 Protease Gene from
RT Clinical Strains of the Hepatitis C Virus."
RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF369235; AAK34560.1; -;
DR InterPro; IPR004109; HCV_NS3.
DR Pfam; PFC2907; HCV_NS3; 1.
KW Protease.
FT NON_TER 1
FT NON_TER 181
SQ SEQUENCE 181 AA; 19130 MW; 85D91969299B7C35 CRC64;
Query Match 99.2%; Score 945; DB 12; Length 181;
Best Local Similarity 99.4%; Pred No. 1.9e-88;
Matches 180; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 APITAYAQTRGLGCIITSLTGRDKNOVEGEVQIVSTAQTFLAICINGVCWTYVHCAG 61
|||||
DB 1 APITAYAQTRGLGCIITSLTGRDKNOVEGEVQIVSTAQTFLAICINGVCWTYVHCAG 60
QY 62 TTTIASPGVPIQMYTNVDKDLGVHPAPQGSRSLSLTPCTCGSSDLYLVTRHADVIPVRRRG 121
|||||
DB 61 TTTIASPGVPIQMYTNVDKDLGVHPAPQGSRSLSLTPCTCGSSDLYLVTRHADVIPVRRRG 120

QY 122 DSRGSLSPRPISVYKSGSGPILCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 181
DB DSRGSLSPRPISVYKSGSGPILCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
QY 182 S 182
DB 181 S 181

RESULT 2

Q91FE5 PRELIMINARY; PRI: 181 AA.
ID Q91FE5
AC Q91FE5
DT 01-DEC-2001 (TrEMBLrel. 15, Created)
DI 01-DEC-2001 (TrEMBLrel. 15, Last sequence update)
DE NS3 protease (Fragment).
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN:P-4;
RA Holland-Staley C.A., Kovari L.C., Giesberg E., Meyers D.L.;
RT "Genetic Diversity and Response to IFN of the NS3 Protease Gene from
Clinical Strains of the Hepatitis C Virus."
RL Submitted (APR-2001) to the EMBL/GenBank/DDBJ databases.
DR EMBL: AF169218; AA:54543.1;
DI InterPro: IPR004139; HCV_NS3.
DR Pfam: PF02907; HCV_NS3; 1.
KW Protease.
FT NON_TER 181 181
SQ SEQUENCE 181 AA: 19-130 MW: 85D9.864294B3C35 CRC64;

Query Match 99.2%; Score 945; DA 12; Length 182;

Best Local Similarity 99.4%; Pred. No. 1.9e-88;

Matches 180; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 APIYAAQOTRGLGCGITSLTGDRKNQVEGVQIVSTAAQTFLATCINGVCTVYHGA 61
DB 1 APIYAAQOTRGLGCGITSLTGDRKNQVEGVQIVSTAAQTFLATCINGVCTVYHGA 60
QY 62 TRIASPKGPIQIYQVYVNDKDLVGNWPAQGSRSITPCTCGSSDLYLVTHADVIPVRR 121
DB 61 TRIASPKGPIQIYQVYVNDKDLVGNWPAQGSRSITPCTCGSSDLYLVTHADVIPVRR 120
QY 122 DSRGSLSPRPISVYKSGSGPILCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 181
DB 121 DSRGSLSPRPISVYKSGSGPILCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
QY 182 S 182
DB 181 S 181

RESULT 3

Q81756 PRELIMINARY; PRI: 2436 AA.
ID Q81756
AC Q81756
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DI 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DE Genome polyprotein (Fragment).
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RA Choo Q.-L., Richman K., Han J.;
RT "The nucleotide sequence of the Hepatitis C viral genome."

Submitted (MAY-1990) to the EMBL/GenBank/DDBJ databases.

RL EMBL: M32084; AA:45677.1;
DR HSP: P27958; IAIY
DR InterPro: IPR001410; DEAD.
DR InterPro: IPR002551; HCV_NS1.
DR InterPro: IPR002518; HCV_NS2.
DR InterPro: IPR004109; HCV_NS3.
DR InterPro: IPR000745; HCV_NS4a.
DR InterPro: IPR001490; HCV_NS4b.
DR InterPro: IPR002868; HCV_NS5a.
DR InterPro: IPR002166; HCV_RDRP.
DR InterPro: IPR001650; Helicase_C.
DR InterPro: IPR007095; RNA_po_DS_PS.
DR InterPro: IPR007094; RNA_po_PSVir.
DR Pfam: PF01560; HCV_NS1; 1.
DR Pfam: PF01518; HCV_NS2; 1.
DR Pfam: PF02907; HCV_NS3; 1.
DR Pfam: PF01006; HCV_NS4a; 1.
DR Pfam: PF01001; HCV_NS4b; 1.
DR Pfam: PF01506; HCV_NS5a; 1.
DR Pfam: PF00271; Helicase_C; 1.
DR Pfam: PF00598; Viral_RDRP; 1.
DR Pfam: PF0186062; HCV_NS1; 1.
DR PRODOM: PD186062; HCV_NS1; 1.
DR SMART: SK03487; RDRP_POSITIVE; 1.
DR PROSITE: PS50507; RDRP_VIRAL; 1.
DR PROSITE: PS50521; RDRP_VIRAL; 1.
KW ATP-binding; Coat protein; Envelope protein; Glycoprotein; Helicase;
KW Hydrolase; Nonstructural protein; Polyprotein;
KW RNA-directed RNA polymerase; Transferase; Transmembrane.
FT NON_TER 2436 2436
SQ SEQUENCE 2436 AA: 264734 MW: L7B9872900BE3125 CRC64;

Query Match 99.0%; Score 943; DA 12; Length 2436;

Best Local Similarity 98.4%; Pred. No. 7.1e-87;

Matches 279; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 MAPITAYAAQOTRGLGCGITSLTGDRKNQVEGVQIVSTAAQTFLATCINGVCTVYHGA 60
DB 576 LAPITAYAAQOTRGLGCGITSLTGDRKNQVEGVQIVSTAAQTFLATCINGVCTVYHGA 635
QY 61 GTETASPKGPIQIYQVYVNDKDLVGNWPAQGSRSITPCTCGSSDLYLVTHADVIPVRR 120
DB 636 GRTIASPKGPIQIYQVYVNDKDLVGNWPAQGSRSITPCTCGSSDLYLVTHADVIPVRR 695
QY 121 GDSRGSLLSPRPISVYKSGSGPILCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
DB 696 GDSRGSLLSPRPISVYKSGSGPILCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 755
QY 181 RS 182
DB 756 RS 757

RESULT 4

Q91FE5 PRELIMINARY; PRI: 3011 AA.
ID Q91FE5
AC Q91FE5
DI 01-OCT-2000 (TrEMBLrel. 15, Created)
DI 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Genome polyprotein.
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=21262212; PubMed=11369872;
RA Lanford R.E., Lee H., Chavez D., Guerra B., Brasky K.M.;
RT "Infectious cDNA clone of the hepatitis C virus genotype 1 prototype
sequence."
RL J. Gen. Virol. 82:1291-1297(2001).


```

RP SEQUENCE FROM N.A.
RC STRAIN=HC-J1;
RA Okamoto H.;
RL Submitted (DEC-1992) to the EMBL/GenBank/DBJ databases.
RN [5]
RP SEQUENCE FROM N.A.
RC STRAIN=HC-J1;
RX MEDLINE=94174722; PubMed=7510436;
RA Mink M., Benichou S., Madaule P., Liollais P., Prince A.,
RA Inchausti G.;
RI "Characterization and mapping of a B-cell immunogenic domain in
RT hepatitis C virus E2 glycoprotein using a yeast peptide library.";
RL Virology 200;246-255(1994).
CC !- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
CC LIPID PROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS: A
CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
CC PROTEIN C AND MRNA (BY SIMILARITY).
CC EMBL: D10749; BAA01582.1; -.
DR HSP: P27958; 1HE1.
DR InterPro: IPR001410; DEAD3.
DR InterPro: IPR002522; HCV_capsid.
DR InterPro: IPR002521; HCV_core.
DR InterPro: IPR002519; HCV_env.
DR InterPro: IPR002531; HCV_NS2.
DR InterPro: IPR002518; HCV_NS2.
DR InterPro: IPR004109; HCV_NS3.
DR InterPro: IPR000745; HCV_NS4a.
DR InterPro: IPR001490; HCV_NS4b.
DR InterPro: IPR002868; HCV_NS5a.
DR InterPro: IPR002166; HCV_NS5a.
DR InterPro: IPR001650; Helicase_C.
DR InterPro: IPR007095; RNA_pol_DS_Ps.
DR InterPro: IPR007094; RNA_pol_PSVir.
DR Pfam: PF01543; HCV_capsid; 1.
DR Pfam: PF01542; HCV_core; 1.
DR Pfam: PF01539; HCV_env; 1.
DR Pfam: PF01560; HCV_NS1; 1.
DR Pfam: PF01538; HCV_NS2; 1.
DR Pfam: PF02907; HCV_NS3; 1.
DR Pfam: PF01005; HCV_NS4a; 1.
DR Pfam: PF01001; HCV_NS4b; 1.
DR Pfam: PF01506; HCV_NS5a; 1.
DR Pfam: PF00271; Helicase_C; 1.
DR Pfam: PF00598; Viral_Rdrp; 1.
DR ProDom: PD186062; HCV_NS1; 1.
DR SMART: SM00487; DEXDC; 1.
DR PROSITE: PS00407; RDRP_POSITIVE; 1.
DR PROSITE: PS0521; RDRP_VIRAL; 1.
KW ATP-binding; Coat protein; Envelope protein; Glycoprotein; Helicase;
KW Hydrolase; Nonstructural protein; Polyprotein;
KW RNA-directed RNA polymerase; Transferase; Transmembrane.
SQ SEQUENCE 3011 AA: 327112 MW: 97905202504535 CRC64;

Query Match 98.5%; Score 930; DB 12; Length 3011;
Best Local Similarity 97.8%; Pred. No. 2.4e-86;
Matches 178; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 1 MAPITAYAAQQTGRLGCGIITSITGRDNKNQVEGEVQIVSTAAQTFLATCINGVCMVYVHGA 60
DB 1026 LAPITAYAAQQTGRLGCGIITSITGRDNKNQVEGEVQIVSTAAQTFLATCINGVCMVYVHGA 1085

QY 61 GRTIASPKGPVIOYNTVDKLVGWPAQPGSRSLTPCTCGSSDLYLVTRHADVTPVRRR 120
DB 1086 GRTIASPKGPVIOYNTVDKLVGWPAQPGSRSLTPCTCGSSDLYLVTRHADVTPVRRR 1145

QY 121 GDSRGSLLSPRPISYIKGSSGGPLLCAGHAGVIFRAAVCTRGVAKAVDTPVSELETTM 160
DB 1146 GDSRGSLLSPRPISYIKGSSGGPLLCAGHAGVIFRAAVCTRGVAKAVDTPVSELETTM 1255

QY 181 RS 162
DB 1206 RS 1207

```

```

RESULT 13
Q91RS4 PRELIMINARY: PRT; 181 AA.
AC Q91RS4;
DT 31-DEC-2001 (TrEMBLrel. 19, Created)
DT 31-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 31-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE NS3 protease (Fragment).
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=PC.23;
RA Holland-Staley C.A., Kovari L.C., Golenberg E., Mayers D.L.;
RT "Genetic Diversity and response to IFN of the NS3 Protease Gene from
RT Clinical Strains of the Hepatitis C Virus.";
RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF369219; AAK54544.1; -.
DR InterPro: IPR004109; HCV_NS3.
DR Pfam: PF02907; HCV_NS3; 1.
DR Protease.
KW NON_TER 1 1
FT NON_TER 181 181
SQ SEQUENCE 181 AA: 19059 MW: 1E5347AE8B7E5C9 CRC64;

Query Match 98.4%; Score 938; DB 12; Length 181;
Best Local Similarity 98.3%; Pred. No. 9.6e-88;
Matches 178; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 2 APITAYAAQQTGRLGCGIITSITGRDNKNQVEGEVQIVSTAAQTFLATCINGVCMVYVHGA 61
DB 1 APITAYAAQQTGRLGCGIITSITGRDNKNQVEGEVQIVSTAAQTFLATCINGVCMVYVHGA 60

QY 62 TRTIASPKGPVIOYNTVDKLVGWPAQPGSRSLTPCTCGSSDLYLVTRHADVTPVRRR 121
DB 61 TKTIASPKGPVIOYNTVDKLVGWPAQPGSRSLTPCTCGSSDLYLVTRHADVTPVRRR 120

QY 122 DSRGSLSPRPISYIKGSSGGPLLCAGHAGVIFRAAVCTRGVAKAVDTPVSELETTM 181
DB 121 DSRGSLSPRPISYIKGSSGGPLLCAGHAGVIFRAAVCTRGVAKAVDTPVSELETTM 180

QY 182 S 162
DB 181 S 161

RESULT 14
Q91RS8 PRELIMINARY: PRT; 181 AA.
AC Q91RS8;
DT 01-DEC-2001 (TrEMBLrel. 19, Created)
DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE NS3 protease (Fragment).
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=PC.176;
RA Holland-Staley C.A., Kovari L.C., Golenberg E., Mayers D.L.;
RT "Genetic Diversity and response to IFN of the NS3 Protease Gene from
RT Clinical Strains of the Hepatitis C Virus.";
RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF369225; AAK54550.1; -.
DR InterPro: IPR004109; HCV_NS3.
DR Pfam: PF02907; HCV_NS3; 1.
DR Protease.
KW NON_TER 1 1
FT NON_TER 181 181

```

Job time : 109 secs

FT NON_TER 181 181
SQ SEQUENCE 181 AA: 19114 MW: 574AC47AE8AEFFD2 CRC64:

Query Match 98.4%; Score 938; DB 12; Length 181;
Best Local Similarity 99.3%; Pred. No. 9.6e-88;
Matches 178; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 2 APTIYAQQTGRLGGCGIITSLTGKKNQVEVEVYSTANQTECATCINGVGVTVYHAG 61
DB 1 APTIYAQQTGRLGGCGIITSLTGKKNQVEVEVYSTANQTECATCINGVGVTVYHAG 60
QY 62 TRTIASPKGPVIQMYTNVCKDLVWKPAPQKSRSLTFCGSSDLYLVTRHADVTPVRRG 121
DB 61 TRTIASPKGPVIQMYTNVCKDLVWKPAPQKSRSLTFCGSSDLYLVTRHADVTPVRRG 120
QY 122 DSRGSLSPRPISYLGKSSGGPLCPACAHVGFRAAVCTRGVAKAVDFIPVSLPTTMR 181
DB 121 DSRGSLSPRPISYLGKSSGGPLCPACAHVGFRAAVCTRGVAKAVDFIPVSLPTTMR 180
QY 182 S 182
DB 181 S 181

RESULT 15
Q91RT3 PRELIMINARY; PRI: 181 AA.
AC Q91RT3;
DT 01-DEC-2001 (TrEMBLrel. 19, Created)
DI 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE NS3 protease (Fragment).
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Pt.11;
RA Holland-Staley C.A., Kovari L.C., Golenberg E., Kayors D.L.;
RT "Genetic Diversity and Response to IFN of the NS3 Protease Gene from
Clinical Strains of the Hepatitis C Virus."
RL Submitted (AFR-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF369220; AAK54545.1;
DR InterPro: IPR004109; HCV_NS3;
DR Pfam: PF02907; HCV_NS3; 1.
KW Protease.
KW NON_TER 1 1
FT NON_TER 181 181
SQ SEQUENCE 181 AA: 19116 MW: 6648807F49B1D33 CRC64:

Query Match 98.4%; Score 938; DB 12; Length 181;
Best Local Similarity 98.3%; Pred. No. 4.6e-85;
Matches 178; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 2 APTIYAQQTGRLGGCGIITSLTGKKNQVEVEVYSTANQTECATCINGVGVTVYHAG 61
DB 1 APTIYAQQTGRLGGCGIITSLTGKKNQVEVEVYSTANQTECATCINGVGVTVYHAG 60
QY 62 TRTIASPKGPVIQMYTNVCKDLVWKPAPQKSRSLTFCGSSDLYLVTRHADVTPVRRG 121
DB 61 TRTIASPKGPVIQMYTNVCKDLVWKPAPQKSRSLTFCGSSDLYLVTRHADVTPVRRG 120
QY 122 DSRGSLSPRPISYLGKSSGGPLCPACAHVGFRAAVCTRGVAKAVDFIPVSLPTTMR 181
DB 121 DSRGSLSPRPISYLGKSSGGPLCPACAHVGFRAAVCTRGVAKAVDFIPVSLPTTMR 180
QY 182 S 182
DB 181 S 181

GenCore version 5.1.6
Copyright (c) 1993 - 2003 Computer Ltd.

OM protein - protein search, using sw model

Run on: September 26, 2003, 22:20:58.3 Search time 72 seconds

(without alignments)
370,362 Million cell updates/sec

Title: US-09-965-594-1

Perfect score: 953

Sequence: 1 MAFITAYACQTRGLGCLIT.....GVAKAVGEIFVRSLEETWMS 182

Scoring table: BLOSUM62

Gapop 10.0, Gapext 0.5

Searched: 1107863 seqs, 158726573 residues

Total number of hits satisfying chosen parameters: 1107862

Minimum DB seq length: 0

Maximum DB seq length: 2000300000

Post-processing: Minimum Match 94

Listing first 45 summaries

Database: A_Geneseq_19Jun03.*

1: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1980.DAT.*
2: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1981.DAT.*
3: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1982.DAT.*
4: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1983.DAT.*
5: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1984.DAT.*
6: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1985.DAT.*
7: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1986.DAT.*
8: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1987.DAT.*
9: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1988.DAT.*
10: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1989.DAT.*
11: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1990.DAT.*
12: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1991.DAT.*
13: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1992.DAT.*
14: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1993.DAT.*
15: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1994.DAT.*
16: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1995.DAT.*
17: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1996.DAT.*
18: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1997.DAT.*
19: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1998.DAT.*
20: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA1999.DAT.*
21: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA2000.DAT.*
22: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA2001.DAT.*
23: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA2002.DAT.*
24: /SIDSL/qcgdata/geneseq/geneseq-emb1/AA2003.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	946	99.3	586	AAE18689	HCV-1 NS3/4a mutant
2	946	99.3	586	AAE18689	Hepatitis C virus
3	946	99.3	586	AAE18689	HCV-1 NS3/4a confo
4	944	99.1	3011	AAE18689	HCV genomic amino
5	943	99.0	609	AAE18689	Hepatitis C virus
6	943	99.0	1766	AAE18689	Sequence encoded i
7	943	99.0	1766	AAE18689	Protein sequence o
8	943	99.0	2261	AAE18689	Peptide encoded by
9	943	99.0	2301	AAE18689	Sequence encoded i

10	943	99.0	2436	10	AAE18689	Sequence encoded i
11	943	99.0	2436	10	AAE18689	Peptide encoded by
12	943	99.0	2772	21	AAE18689	Protein encoded by
13	943	99.0	2816	14	AAE18689	HCV-1 polyprotein.
14	943	99.0	2894	16	AAE18689	Composite hepatiti
15	943	99.0	2955	20	AAE18689	Amino acid sequenc
16	943	99.0	2955	21	AAE18689	Polyprotein encode
17	943	99.0	3011	13	AAE18689	Compiled HCV sequ
18	943	99.0	3011	14	AAE18689	Hepatitis C virus
19	943	99.0	3011	17	AAE18689	Hepatitis C virus
20	943	99.0	3011	18	AAE18689	HCV polyprotein.
21	943	99.0	3011	19	AAE18689	HCV polyprotein.
22	943	99.0	3011	23	AAE18689	Hepatitis C virus
23	943	99.0	3011	23	AAE18689	Hepatitis C virus
24	942	98.8	632	23	AAE18689	HCV polyprotein la
25	942	98.8	632	23	AAE18689	Hepatitis C virus
26	942	98.8	686	23	AAE18689	Hepatitis C virus
27	942	98.8	686	23	AAE18689	Hepatitis C virus
28	942	98.8	686	23	AAE18689	Hepatitis C virus
29	942	98.8	686	23	AAE18689	Hepatitis C virus
30	942	98.8	686	23	AAE18689	Hepatitis C virus
31	942	98.8	686	23	AAE18689	Hepatitis C virus
32	942	98.8	686	23	AAE18689	Hepatitis C virus
33	942	98.8	686	23	AAE18689	Hepatitis C virus
34	942	98.8	686	23	AAE18689	Hepatitis C virus
35	942	98.8	686	23	AAE18689	Hepatitis C virus
36	942	98.8	686	23	AAE18689	Hepatitis C virus
37	942	98.8	686	23	AAE18689	Hepatitis C virus
38	942	98.8	686	23	AAE18689	Hepatitis C virus
39	942	98.8	686	23	AAE18689	Hepatitis C virus
40	942	98.8	686	23	AAE18689	Hepatitis C virus
41	942	98.8	686	23	AAE18689	Hepatitis C virus
42	942	98.8	686	23	AAE18689	Hepatitis C virus
43	942	98.8	686	23	AAE18689	Hepatitis C virus
44	942	98.8	686	23	AAE18689	Hepatitis C virus
45	942	98.8	686	23	AAE18689	Hepatitis C virus

ALIGNMENTS

RESULT 1

AAE18689

10 AAE18689 standard: Protein; 686 AA.

XX AAE18689;

XX 17-MAY-2002 (first entry)

XX HCV-1 NS3/4a mutant conformational antigen.

XX Hepatitis C virus; NS3/4a antigen; HCV infection; mutant; muten.

XX Hepatitis C virus type 1.

XX Synthetic.

XX Key

XX Location/Qualifiers

XX Misc-difference 403

XX /note- "Wild type Thr substituted with Pro"

XX Misc-difference 404

XX /note- "Wild type Ser substituted with Ile"

XX WC200196675-A2.

XX 20-DEC-2001.

XX 14-JUN-2001; 2001WO-0519369.

XX 15-JUN-2000; 2000US-212082P.

XX 02-APR-2001; 2001US-280811P.

XX 02-APR-2001; 2001US-280867P.

XX (CHIR) CHIRON CORP.

```

XX Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Coit D;
PI Medina-Selby A;
XX WP1: 2002-179522/23.
DR N-PSDB; AAD2979s.
XX
XX Immunassay solid support useful for detecting hepatitis C virus
XX infection in a biological sample, comprises at least one of HCV
XX anti-core antibody and HCV NS3/4a epitope, bound to the support.
XX
XX Example 2: Fig 4: 87pp: English.
XX
XX The present invention relates to hepatitis C virus (HCV) core antigen
XX and NS (nonstructural) 3/4a antibody combination assay that can detect
XX both HCV antigens and antibodies present in a sample using a single
XX solid matrix as well as immunoassay solid supports for use in the assay.
XX The solid support is useful for detecting HCV infection in a biological
XX sample. The present sequence is HCV-1 NS3/4a mutant conformational
XX antigen. This sequence is used in the exemplification of the invention.
XX
XX Sequence 686 AA:
XX
XX Query Match 99.3%; Score 946; DA 23; Length 686;
XX Best Local Similarity 98.9%; Pred. No. 1.2e-89;
XX Matches 180; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 MAPITAYAQOTRGLGCIITSLTGKDKNQVEGEVQVSTAAQTFLATCINGVCTWYVHGA 60
XX DB 1 MAPITAYAQOTRGLGCIITSLTGKDKNQVEGEVQVSTAAQTFLATCINGVCTWYVHGA 60
XX
XX QY 61 GTRIASPQGPVIOMYTNVDKDLVGNWPAPOGSRSLTPTCGSSDLVLTTHADVIPVRRR 120
XX DB 61 GTRIASPQGPVIOMYTNVDKDLVGNWPAPOGSRSLTPTCGSSDLVLTTHADVIPVRRR 120
XX
XX QY 121 GDSRGSLLSPRPISYLGKSGGGLPCPAGHAVGIFRAAVCTRCGVAKAVDFIPVESLETH 180
XX DB 121 GDSRGSLLSPRPISYLGKSGGGLPCPAGHAVGIFRAAVCTRCGVAKAVDFIPVESLETH 180
XX
XX QY 181 RS 182
XX DB 181 RS 182
XX
XX RESULT 3
XX ABG72261
XX ID ARG72261 standard; Protein: 686 AA.
XX AC ABG72261;
XX DT 06-MAR-2003 (first entry)
XX DE HCV-1 NS3/4a conformational antigen.
XX

```

```

FR 15-JUN-2000; 2000US-212082P.
FR 02-APR-2001; 2001US-280811P.
FR 02-APR-2001; 2001US-280867P.
XX (CHIR) CHIRON CORP.
XX Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Coit D;
PI Medina-Selby A;
XX WP1: 2002-090228/12.
DR N-PSDB; ABK15344.
XX
XX Immunassay solid support, useful for detecting hepatitis C virus
XX infection in biological sample, comprises HCV NS3/4a conformational
XX epitope and multiple epitope fusion antigen bound to the support.
XX Claim 5; Fig 3: 92pp: English.
XX
XX The present invention relates to a new immunoassay solid support
XX consisting essentially of at least one hepatitis C virus (HCV) NS3/4a
XX conformational epitope and a multiple epitope fusion antigen (MEFA),
XX bound to the support. The NS3/4a conformational epitope and/or
XX MEFA reacts specifically with anti-HCV antibodies present in a biological
XX sample from an HCV-infected individual. The immunoassay of the invention
XX is useful for detecting hepatitis C virus infection in a biological
XX sample. The method of the invention provides a sensitive, accurate
XX diagnostic and prognostic tool to provide adequate patient care and to
XX prevent transmission of HCV by blood and by blood products, or by
XX personal contact. Use of NS3/4a conformational epitope in combination
XX with MEFA, provides a sensitive and reliable method for detecting early
XX HCV seroconversion. Use of MEFA has the added advantages of decreasing
XX masking problems, improving sensitivity in detecting antibodies by
XX allowing a greater number of epitopes on a unit surface area of
XX substrate, and improving substrate. Detection accuracy is increased and
XX the incidence of false results is reduced because of the identification
XX and the use of highly immunogenic HCV antigens which are present during
XX the early stages of HCV seroconversion. The present amino acid sequence
XX represents the non-structural protein NS3/4a conformational epitope of
XX the invention.
XX
XX Sequence 686 AA:
XX
XX Query Match 99.3%; Score 946; DA 23; Length 686;
XX Best Local Similarity 98.9%; Pred. No. 1.2e-89;
XX Matches 180; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 MAPITAYAQOTRGLGCIITSLTGKDKNQVEGEVQVSTAAQTFLATCINGVCTWYVHGA 60
XX DB 1 MAPITAYAQOTRGLGCIITSLTGKDKNQVEGEVQVSTAAQTFLATCINGVCTWYVHGA 60
XX
XX QY 61 GTRIASPQGPVIOMYTNVDKDLVGNWPAPOGSRSLTPTCGSSDLVLTTHADVIPVRRR 120
XX DB 61 GTRIASPQGPVIOMYTNVDKDLVGNWPAPOGSRSLTPTCGSSDLVLTTHADVIPVRRR 120
XX
XX QY 121 GDSRGSLLSPRPISYLGKSGGGLPCPAGHAVGIFRAAVCTRCGVAKAVDFIPVESLETH 180
XX DB 121 GDSRGSLLSPRPISYLGKSGGGLPCPAGHAVGIFRAAVCTRCGVAKAVDFIPVESLETH 180
XX
XX QY 181 RS 182
XX DB 181 RS 182
XX
XX RESULT 3
XX ABG72261
XX ID ARG72261 standard; Protein: 686 AA.
XX AC ABG72261;
XX DT 06-MAR-2003 (first entry)
XX DE HCV-1 NS3/4a conformational antigen.
XX

```

KW Immunobassay solid support; Hepatitis C Virus type-1; HCV-1;
 KW NS3/4a conformational epitope; multiple epitope fusion antigen;
 KW MEFA; anti-HCV antibody; NS3/4a conformational antigen;
 KW HCV infection; mutant; mutagen.
 OS
 OS Hepatitis C virus type 1.
 OS Synthetic.
 XX
 XX Key Location/Qualifiers
 FH 2...586
 FT Region /note: "Corresponds to amino acid residues 1027-1711
 FT of HCV-1 NS3/4a polypeptide"
 FT Misc-difference 403
 FT /note: "Substitution of wild-type Thr to Pro"
 FT Misc-difference 404
 FT /note: "Substitution of wild-type Ser to I-Le"
 XX
 XX US2002146685-A1.
 XX
 XX 10-OCT-2002.
 XX
 XX 14-JUN-2001; 2001US-0861654.
 XX
 XX 15-JUN-2000; 2000US-212082P.
 PR 02-APR-2001; 2001US-280811P.
 PR 02-APR-2001; 2001US-280857P.
 XX
 XX (CHIEF) CHIEN D Y.
 PA (ARCA) ARCANDEL P.
 PA (TAND) TANDESKE L.
 PA (GEOR) GEORGE-NASCIMENTO C.
 PA (COLT) COLT D.
 PA (MEDT) MEDINA-SELBY A.
 XX
 XX Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Colt D;
 PI Medina-Selby A;
 XX WPI: 2003-147573/14.
 XX N-PSDB; ARX14410.
 DR
 XX Immunobassay solid support for detecting Hepatitis C Virus infection in
 PT biological samples, comprises Hepatitis C Virus conformational epitope
 PT and multiple epitope fusion antigen.
 XX
 XX Claim 2: Fig 3A-3D; 45pp; English.
 PS
 XX The present invention relates to immunoassays comprising Hepatitis C
 CC virus (HCV) NS3/4a conformational epitope and multiple epitope fusion
 CC antigen (MEFA), bound to a solid support. The NS3/4a epitope and/or
 CC the multiple epitope fusion antigen react with anti-HCV antibodies
 CC present in a biological sample from an HCV-infected individual. The
 CC immunoassays and methods of the invention are useful for detecting
 CC HCV infection in a biological sample. The inventive immunoassay solid
 CC support provides a sensitive and reliable method for detecting early
 CC HCV seroconversion. The assays can detect HCV infection caused by any
 CC six known genotypes of HCV. The use of the multiple epitope fusion
 CC proteins decreases masking problems, improves sensitivity in detecting
 CC antibodies by allowing a greater number of epitopes on a unit area
 CC of substrate, and improves selectivity. The present sequence
 CC represents HCV type 1 (HCV-1) NS3/4a conformational antigen, a mutant
 CC of the HCV-1 NS3/4a polypeptide.
 XX
 XX Sequence 586 AA:
 SQ
 Query Match 99.3%; Score 346; DR 24; Length 686;
 Best Local Similarity 98.9%; Pred. No. 1.2e-89;
 Matches 180; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MAPITAAQQTGRLGCLITSLTGRKNOVEGVQIVSTAAQTFLATCINGVCTVYHGA 60
 DB 1 MAPITAAQQTGRLGCLITSLTGRKNOVEGVQIVSTAAQTFLATCINGVCTVYHGA 60
 QY 61 GTRTIASPKGPVQMTYNVDKLVGWPAPOGGRSITPTCGSSDLYLVTRHADVIPVRRR 120

DB 61 GTRTIASPKGPVQMTYNVDKLVGWPAPOGGRSITPTCGSSDLYLVTRHADVIPVRRR 120
 QY 121 GDSKGSLLSPRPISYLGKSSGGPPLCPAGHAGVCFRAAVCTIRGVAKAVDFIPVESLETMM 180
 DB 121 GDSKGSLLSPRPISYLGKSSGGPPLCPAGHAGVCFRAAVCTIRGVAKAVDFIPVENLETTM 180
 QY 181 RS 182
 DB 181 RS 182
 RESULT 4
 AAR4C120
 ID AAR4C120 standard; Protein: 3011 AA.
 AC AAR4C120;
 XX 25-MAR-2003 (updated)
 DT 27-JAN-1994 (first entry)
 XX
 XX HCV genomic amino acid sequence isolated from infected human LG.
 XX
 XX Hepatitis C Virus; Non-A, non-B Hepatitis Virus; HCV; NANBHV;
 KW human growth hormone; HGH; secretion signal; fusion protein;
 KW vaccine.
 XX
 XX Hepatitis C Virus.
 OS WO9315193-A1.
 XX
 XX 05-AUG-1993.
 PD
 XX 29-JAN-1993; 93WO-US00907.
 PF
 XX 31-JAN-1992; 92US-0630C24.
 PR
 XX (ABBO) ABBOTT LAB.
 PA
 XX Bode SL, Casey JM, Desai SM, Devare SG, Frail DE;
 PT Yamaguchi J, Zeck BJ;
 XX WPI: 1993-258673/32.
 DR
 XX New plasmid pHCV-162 is a mammalian expression systems for HCV
 PT proteins - useful for diagnosing HCV infection and as vaccines
 PT for preventing HCV infection
 PS
 XX Example 1; Page 39-49; 100pp; English.
 CC
 CC RNA was isolated from the plasma of a HCV seropositive human
 CC (designated "IG") and cDNA was prepared from it. The cDNA was
 CC PCR amplified using specific primers with sequences based
 CC on the prototype HCV-1 cDNA sequence (GENBANK M62321). Further
 CC amplification using nested primers resulted in 7 adjacent HCV DNA
 CC fragments which could be assembled into a full-length sequence. The
 CC DNA sequence was determined and translated into the genomic amino
 CC acid sequence. Comparison of the LG genomic amino acid sequence
 CC with that from HCV-1 showed 134 amino acid differences.
 CC (Updated on 25-MAR-2003 to correct PN field.)
 XX
 XX Sequence 3011 AA:
 SQ
 Query Match 99.1%; Score 944; DR 14; Length 3011;
 Best Local Similarity 98.4%; Pred. No. 1.4e-88;
 Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MAPITAAQQTGRLGCLITSLTGRKNOVEGVQIVSTAAQTFLATCINGVCTVYHGA 60
 DB 1026 LAPITAAQQTGRLGCLITSLTGRKNOVEGVQIVSTAAQTFLATCINGVCTVYHGA 1085
 QY 61 GTRTIASPKGPVQMTYNVDKLVGWPAPOGGRSITPTCGSSDLYLVTRHADVIPVRRR 120


```

DE 12f through 15e.
XX Hepatitis C virus (HCV): non-A, non-B hepatitis (HANBH).
XX Hepatitis C virus.
XX Hepatitis C virus.
XX EP318216-A.
XX 31-MAY-1989.
XX 18-NOV-1988; 88EP-0310922.
XX 18-NOV-1987; 87US-0122714.
XX 30-DEC-1987; 87US-0139886.
XX 26-FEB-1988; 88US-0161072.
XX 06-MAY-1988; 88US-0191263.
XX 26-OCT-1988; 88US-0263584.
XX 14-NOV-1988; 88US-0271450.
XX (CHIR ) CHIRON CORP.
XX Houghton M, Choo QL, Kuo G;
XX WPI; 1989-159274/22.
XX N-PSDB; AAN92103.
XX Purified hepatitis C virus
XX - and associated nucleic acids and polypeptide(s)
XX Claim 13; Figure 32-1 - 32-7; 139 pp; English.
XX It is the sequence encoded in the open reading frame of hepatitis C virus
XX (HCV) cDNA inserts in clones 12f through 15e. It is antigenic and could
XX be used in immunoassay reagents and vaccines and to generate antibodies
XX useful in diagnosis and passive immunotherapy for HCV infection/non-A,
XX non-B hepatitis.
XX (Updated on 25-MAR-2003 to correct PR field.)
XX (Updated on 25-MAR-2003 to correct PI field.)
XX Sequence 2301 AA;
Query Match 99.0%; Score 943; DB 10; Length 2301;
Best Local Similarity 98.4%; Pred. No. 1,2e-88;
Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
QY 1 MAPITAYAOOTRGLGCIITSLTGRDKNQVEGEVCIYSTAAQTFLATCINGVCMVYHGA 60
DE 401 LAPITAYAOOTRGLGCIITSLTGRDKNQVEGEVCIYSTAAQTFLATCINGVCMVYHGA 460
QY 51 GTRTIASPKGPVIOQMTNVKDLVGNWPAQGSRLTPTCGSSDLYLVTRHADVIPVRRR 120
DB 461 GTRTIASPKGPVIOQMTNVDDLVGNWPAQGSRLTPTCGSSDLYLVTRHADVIPVRRR 520
QY 121 GDSKGSLLSPRPISYLVKGSNGGGLPCAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
DB 521 GDSKGSLLSPRPISYLVKGSNGGGLPCAGHAGVIFRAAVCTRGVAKAVDFIPVENLETTM 580
QY 181 RS 182
DB 561 RS 582
RESULT 11
AAP92050
ID AAP92050 standard; protein: 2436 AA.
XX AAP92050;
XX AC
XX 25-MAR-2003 (updated)
XX 02-MAR-1990 (first entry)
XX Sequence encoded in the hepatitis C virus (HCV) cDNA inserts in clones
DE K9-1 through 15e.

```

```

XX Hepatitis C virus (HCV): non-A, non-B hepatitis (HANBH).
XX Hepatitis C virus.
XX EP318216-A.
XX 31-MAY-1989.
XX 18-NOV-1988; 88EP-0310922.
XX 18-NOV-1987; 87US-0122714.
XX 30-DEC-1987; 87US-0139886.
XX 26-FEB-1988; 88US-0161072.
XX 06-MAY-1988; 88US-0191263.
XX 26-OCT-1988; 88US-0263584.
XX 14-NOV-1988; 88US-0271450.
XX (CHIR ) CHIRON CORP.
XX Houghton M, Choo QL, Kuo G;
XX WPI; 1989-159274/22.
XX N-PSDB; AAN92106.
XX Purified hepatitis C virus
XX - and associated nucleic acids and polypeptide(s)
XX Claim 13; Figure 47-1 - 47-8; 139 pp; English.
XX It is the sequence encoded in the open reading frame of hepatitis C virus
XX (HCV) cDNA inserts in clones K9-1 through 15e. It is antigenic and could
XX be used in immunoassay reagents and vaccines and to generate antibodies
XX useful in diagnosis and passive immunotherapy for HCV infection/non-A,
XX non-B hepatitis.
XX (Updated on 25-MAR-2003 to correct PR field.)
XX (Updated on 25-MAR-2003 to correct PI field.)
XX Sequence 2436 AA;
Query Match 99.0%; Score 943; DB 10; Length 2436;
Best Local Similarity 98.4%; Pred. No. 1,3e-88;
Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
QY 1 MAPITAYAOOTRGLGCIITSLTGRDKNQVEGEVCIYSTAAQTFLATCINGVCMVYHGA 60
DB 576 LAPITAYAOOTRGLGCIITSLTGRDKNQVEGEVCIYSTAAQTFLATCINGVCMVYHGA 635
QY 51 GTRTIASPKGPVIOQMTNVKDLVGNWPAQGSRLTPTCGSSDLYLVTRHADVIPVRRR 120
DB 536 GTRTIASPKGPVIOQMTNVDDLVGNWPAQGSRLTPTCGSSDLYLVTRHADVIPVRRR 695
QY 121 GDSKGSLLSPRPISYLVKGSNGGGLPCAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
DB 696 GDSKGSLLSPRPISYLVKGSNGGGLPCAGHAGVIFRAAVCTRGVAKAVDFIPVENLETTM 755
QY 181 RS 182
DB 756 RS 757
RESULT 11
AAP90288
ID AAP90288 standard; protein: 2436 AA.
XX AAP90288;
XX AC
XX 25-MAR-2003 (updated)
XX 19-JUL-2001 (updated)
XX 01-NOV-1989 (first entry)
XX Peptide encoded by composite hepatitis C cDNA.

```



```

KW Hepatitis C virus; clone 15c; clone k9-1; probe; vaccine.
XX
OS Pan troglodytes.
XX
PN GB2212911-A.
XX
XX 26-JUL-1989.
XX
XX 18-NOV-1988; 583B-0027224.
XX
XX 19-NOV-1987; 87US-0122714.
XX
XX 30-DEC-1987; 87US-0139886.
XX
XX 26-FEB-1986; 88US-0161072.
XX
XX 26-OCT-1988; 86US-0263584.
XX
PA (CHIR ) CHIRON CORPORATION.
XX
PI Houghton M, Choo QL, Kuo G;
XX
DR WPI; 1989-215054/30.
XX
DR N-PSDB; AAN9336.
XX
PT Hepatitis C virus gene - used for prodn. of polynucleotide protein.
PI polypeptide(s) and antibodies for diagnosis, prevention and
PT treatment of infection.
XX
PS Disclosure; fig 47-1 to 47-8; 235pp; English.
XX
CC The sequence is the peptide encoded by the composite hepatitis C
CC virus (HCV) cDNA of AAN9336. The polypeptides are used to
CC diagnose HCV-induced NANBH. To raise antibodies for
CC immunoassay or treatment, or to produce vaccines.
CC (N.B. This record was resubmitted to correct errors in the sequence.)
XX (Updated on 25-MAR-2003 to correct PR field.)
XX
SQ Sequence 2436 AA;
Query Match 99.0%; Score 943; DB 10; Length 2436;
Best Local Similarity 98.4%; Prod. No. 1.3e-86;
Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
QY I MAPITAYAOQTGRLGCIITSLTGRDKNOVEGEVQIVSTAAQTFLATCINGVCTVYHGA 60
Db :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
576 LAPITAYAOQTGRLGCIITSLTGRDKNOVEGEVQIVSTAAQTFLATCINGVCTVYHGA 635
QY 61 GTTITASPKGPVIOMYTNVDKLVGWPAPOGSRSITPCTCGSSDLYLVTRHADVIPVRRR 120
Db :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
636 GTTITASPKGPVIOMYTNVDQDLVGVWPAPOGSRSITPCTCGSSDLYLVTRHADVIPVRRR 695
QY 121 GDSRGLSPRPISYLGSGGSPILCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
Db :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
696 GDSRGLSPRPISYLGSGGSPILCPAGHAGVIFRAAVCTRGVAKAVDFIPVENLETTM 755
QY 181 RS 182
Db :||
756 RS 757
RESULT 12
AAB18540
ID AAB18540 standard; Protein: 2772 AA.
XX
XX AAB18540;
XX
XX 15-JAN-2001 (first entry)
XX
XX Protein encoded by a cDNA compiled Hepatitis C virus cDNA clones.
XX
XX Hepatitis C virus; HCV; antisense polynucleotide; polypeptide;
XX viral infectivity; viral replication.
XX
XX Hepatitis C virus.
XX

```

```

PN EP1034785-A2.
XX
XX 13-SEP-2000.
XX
XX 16-MAR-1990; 2000EP-0109602.
XX
XX 17-MAR-1989; 89US-0325338.
XX
XX 20-APR-1989; 89US-0341334.
XX
XX 18-MAY-1989; 89US-0355002.
XX
XX 16-MAR-1990; 90EP-0302866.
XX
PA (CHIR ) CHIRON CORP.
XX
PI Houghton M, Choo Q, Kuo G;
XX
DR WPI; 2000-566891/53.
XX
DR N-PSDB; AAN75296.
XX
PT Novel composition comprising a hepatitis C virus antisense
PI polynucleotide which is complementary to or corresponds to a sense
PT strand of the virus genome, and selectively hybridises to it.
XX
PS Example; Fig 16; 75pp; English.
XX
CC The specification describes a pharmaceutical composition which
CC comprises a hepatitis C virus (HCV) antisense polynucleotide. The
CC HCV is characterized by a positive stranded RNA genome which has
CC 40% homology at the polypeptide level to a HCV polyprotein. The
CC antisense polynucleotide binds to cellular polynucleotides which
CC enhance and/or are required for viral infectivity, replicative
CC ability or chronicity. The antisense polynucleotides may also be
CC designed to bind with high specificity, to be of increased stability,
CC to be stable and to have low toxicity. The composition also comprises
CC an agent which causes viral RNA to be inactive. The composition
CC is used for preventing HCV replication in a system. The present
CC sequence is encoded by a novel HCV cDNA sequence, which is used in the
XX course of the invention.
XX
SQ Sequence 2772 AA;
Query Match 99.0%; Score 943; DB 21; Length 2772;
Best Local Similarity 98.4%; Prod. No. 1.5e-88;
Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
QY I MAPITAYAOQTGRLGCIITSLTGRDKNOVEGEVQIVSTAAQTFLATCINGVCTVYHGA 60
Db :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
912 LAPITAYAOQTGRLGCIITSLTGRDKNOVEGEVQIVSTAAQTFLATCINGVCTVYHGA 971
QY 61 GTTITASPKGPVIOMYTNVDKLVGWPAPOGSRSITPCTCGSSDLYLVTRHADVIPVRRR 120
Db :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
972 GTTITASPKGPVIOMYTNVDQDLVGVWPAPOGSRSITPCTCGSSDLYLVTRHADVIPVRRR 1031
QY 121 GDSRGLSPRPISYLGSGGSPILCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
Db :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
1032 GDSRGLSPRPISYLGSGGSPILCPAGHAGVIFRAAVCTRGVAKAVDFIPVENLETTM 1091
QY 181 RS 182
Db :||
1092 RS 1093
RESULT 13
AAR34009
ID AAR34009 standard; Protein: 2816 AA.
XX
XX AAR34009;
XX
XX 25-MAR-2003 (updated)
XX
XX 26-JUL-1993 (first entry)
XX
XX HCV-1 polypeptide.
XX
XX Polymerase chain reaction; PCR; amplify; primer; hepatitis C virus;

```

KW HCV; asymptomatic; chronically infected; epitope; viral isolate;
 KW domain; immunological; cross-reactive; envelope protein; vaccine;
 KW gp53(RVDV)/gp55; hog cholera virus; pestivirus; NS5; flavivirus.

XX Hepatitis C virus.

OS KC9356126-A1.

PN 01-APR-1993.

XX 11-SEP-1992: 92XO-0807683.

XX 13-SEP-1991: 91US-3759575.

XX (CHIR) CHIRON CORP.

XX Houghton M, Weiner AJ;

XX WPI; 1993-117458/14.

XX Immunoreactive hepatitis C virus polypeptide constructs - cont'd.
 PT at least 2 sequences from the first variable domain of distinct
 PT HCV isolates

PS Disclosure; Fig 9; 106pp; English.

XX This sequence represents the entire hepatitis C virus polyprotein.
 CC HCV is a member of the flavivirus family and appears to encode a basic
 CC polypeptide domain ("C") at the N-terminal of the viral polyprotein,
 CC followed by two glycoprotein domains ("E1", "E2/NS1"), upstream of the
 CC nonstructural genes NS2 through NS5. See also AAQ39134-48, AAR35982-
 CC 4008 and AAR3688-89.
 CC (Updated on 25-MAR-2003 to correct PN field.)

XX Query Match 99.0%; Score 943; DS 14; Length 2816;

XX Best Local Similarity 98.4%; Pred. No. 1.7e-88;

XX Matches 179; Conservative 3; Mismatches 9; Indels 0; Gaps 0;

QY 1 MAPTAYAQTRGLGCIITSLTGKKNQVEGEVQIVSTAATGFLATCINGVCTVYHGA 60

Db 1026 LAPITAYAQTRGLGCIITSLTGKKNQVEGEVQIVSTAATGFLATCINGVCTVYHGA 1085

QY 61 GRTIASPKGPVICYMTNVDKDLVGNPAPGSGSLTPTCTGSSDLYLVTRHADVIPVRRR 120

Db 1086 GRTIASPKGPVICYMTNVDKDLVGNPAPGSGSLTPTCTGSSDLYLVTRHADVIPVRRR 1145

QY 121 GDSRGSLSLSPRISYLYKSGSGGPLLCFAGHAGVGFRAAVCTRGVAKAVDFIPVENLETTM 180

Db 1146 GDSRGSLSLSPRISYLYKSGSGGPLLCFAGHAGVGFRAAVCTRGVAKAVDFIPVENLETTM 1205

QY 181 RS 182

Db 1206 RS 1207

RESULT 14

AA70230

XX AAR70230 standard; protein: 2894 AA.

XX AAR70230;

XX 25-MAR-2003 (updated)

DT 07-NOV-1995 (first entry)

XX Composite hepatitis C virus (HC-J1/CDC/CHI).

XX Composite hepatitis C virus; HC-J1/CDC/CHI; HCV; non-A non-B;

KW synthetic antigens; blood screening.

XX Hepatitis C virus.

XX

PN EP644232-A1.

XX 22-MAR-1995.

XX 14-DEC-1990; 94EP-0108611.

XX 14-DEC-1990; 90EP-0124241.

XX 14-DEC-1990; 90EP-0108611.

XX (INNO-) INNOGENETICS NV.

XX Deleys RJ, Maertens G, Pollet D, Van Heuverswyn H;

XX WPI; 1995-116946/16.

XX Synthetic antigens for the detection of hepatitis C virus

PT antibodies - comprise portions of the HCV peptide sequence, for

PT use in screening blood and blood products

XX Disclosure; Fig 1; 51pp; English.

XX AAR70230 is the composite hepatitis C virus (HC-J1/CDC/CHI) protein

CC from which the synthetic HCV antigens described in AAR70210-R70229

CC were derived. These synthetic antigens can be used to screen blood,

CC or blood products for the presence of HCV, they can also be used in

CC various specific assays for the detection of HCV antibodies, and

CC antigens, or as immunogens.

CC (Updated on 25-MAR-2003 to correct PN field.)

CC (Updated on 25-MAR-2003 to correct PF field.)

XX Sequence 2894 AA;

QY Query Match 99.0%; Score 943; DS 16; Length 2894;

Db Best Local Similarity 98.4%; Pred. No. 1.7e-88;

QY Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 MAPTAYAQTRGLGCIITSLTGKKNQVEGEVQIVSTAATGFLATCINGVCTVYHGA 60

Db 1026 LAPITAYAQTRGLGCIITSLTGKKNQVEGEVQIVSTAATGFLATCINGVCTVYHGA 1085

QY 61 GRTIASPKGPVICYMTNVDKDLVGNPAPGSGSLTPTCTGSSDLYLVTRHADVIPVRRR 120

Db 1086 GRTIASPKGPVICYMTNVDKDLVGNPAPGSGSLTPTCTGSSDLYLVTRHADVIPVRRR 1145

QY 121 GDSRGSLSLSPRISYLYKSGSGGPLLCFAGHAGVGFRAAVCTRGVAKAVDFIPVENLETTM 180

Db 1146 GDSRGSLSLSPRISYLYKSGSGGPLLCFAGHAGVGFRAAVCTRGVAKAVDFIPVENLETTM 1205

QY 181 RS 182

Db 1206 RS 1207

RESULT 15

AA714975

XX AAY14975 standard; Protein: 2955 AA.

XX AAY14975;

XX 20-MAR-2003 (updated)

DT 08-NOV-1999 (first entry)

XX Amino acid sequence of HCV-1 ORF.

XX Hepatitis C virus; HCV; J1; J7; HCV-1; non-A, non-B HCV; NANBH;

KW HCV infection; vaccine.

XX Hepatitis C virus.

XX Key Location/Qualifiers

FT Misc-difference 441

FT /note= "encoded by TT"

FT Misc-difference 461

/note- "encoded by CCCC"

FT XX EP339128-A2.
PN XX
XX XX
PD XX 01-SEP-1994.
XX XX
XX XX 17-SEP-1990; 99EP-0101746.
XX XX
XX XX 15-SEP-1989; 89US-0408245.
PR XX 21-DEC-1989; 89US-0456142.
PR XX 17-SEP-1990; 90EP-0310149.
XX XX
XX XX (CHIR) CHIRON CORP.
PA XX (CYAA/) OYA A.
XX XX
PI Cha T. Han J. Houghton M. Irvine BD. Kolberg JA.
PI Miyamura T. Saito I. Weiner AJ.
XX XX
DR WPI: 1999-480843/42.
DR N-PSDB: RA207656.
XX XX
PT New Hepatitis C Virus isolates, useful for diagnosis of Hepatitis
PT Infections and development of vaccines
XX XX
PS Disclosure: Fig 12; 142pf; English.
XX XX
CC The invention provides two new isolates of Hepatitis C virus (HCV), H1
CC and J7. These two isolates comprise nucleotide and amino acid sequences
CC that are distinct from the HCV isolate HCV-1. The nucleotide sequences
CC may be used to detect non-A, non-B HCV (NANBH) polynucleotides by
CC hybridisation for diagnosis of NANBH infections. They may also be used to
CC screen blood donors, donated blood and blood products for this infection.
CC The isolates may also be used to isolate other naturally occurring
CC variants of the virus. The polypeptides may be used as a vaccine for
CC administration to patients to protect against infection with NANBH. The
CC present sequence represents the amino acid sequence of HCV-1 085.
CC (Updated on 20-MAR-2003 to correct pf field.)
CC (Updated on 20-MAR-2003 to correct PR field.)
XX XX
SQ Sequence 2955 AA;

Query Match 99.0%; Score 943; DB 20; Length 2955;
Best Local Similarity 98.4%; Prod. No. 1.7e-98;
Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
QY 1 MAPITAYAOQIRGLLGGIITSLTGRDKNOVGEVOIVSTAAQFLATCINGVGVYVHGA 60
Db 1026 LAPITAYAOQIRGLLGGIITSLTGRDKNOVGEVOIVSTAAQFLATCINGVGVYVHGA 1085
QY 61 GTRITASPKGFVIOYTVNVDQDLVGNFAPQGSRLTFCCTGSSDLYLVTHADVIVPVRK 120
Db 1086 GTRITASPKGFVIOYTVNVDQDLVGNFAPQGSRLTFCCTGSSDLYLVTHADVIVPVRK 1145
QY 121 GDSRGSLLSPRISYLYKSGSGGPLLCPAGHVG:FRAAVTRGVAKAVD:IPVESLETTM 180
Db 1146 GDSRGSLLSPRISYLYKSGSGGPLLCPAGHVG:FRAAVTRGVAKAVD:IPVENLETTM 1205
QY 181 RS 182
Db 1206 RS 1207

Search completed: September 27, 2003, 12:16:53
Job time : 79 secs

FT /label= Thr, Ala
 FT Misc-difference 405
 FT /label= Gln, Pro, Leu
 FT Misc-difference 410
 FT /label= Lys, Arg
 FT Misc-difference 418
 FT /label= Gly, Asp
 FT Misc-difference 430
 FT /label= Asn, Asp
 FT Misc-difference 436
 FT /label= Phe, Leu
 FT Misc-difference 478
 FT /label= Arg, Lys
 FT Misc-difference 759
 FT /label= Leu, Val
 FT Misc-difference 1017
 FT /label= Ser, Asn
 FT Misc-difference 1036
 FT /label= Thr, Ala
 FT Misc-difference 1056
 FT /label= Glu, Asp
 FT Misc-difference 1201
 FT /label= Met, Thr
 FT Misc-difference 1205
 FT /label= Met, Ile
 FT Misc-difference 1255
 FT /label= Asn, Tyr
 FT Misc-difference 1263
 FT /label= Gly, Asp
 FT Misc-difference 1455
 FT /label= Asn, Asp
 FT Misc-difference 1828
 FT /label= Ala, Thr
 FT Misc-difference 1895
 FT /label= Gly, Arg
 FT Misc-difference 1896
 FT /label= Gly, Ile
 FT Misc-difference 2143
 FT /label= Glu, Val
 FT Misc-difference 2144
 FT /label= Asp, Gln
 FT Misc-difference 2462
 FT /label= Cys, Arg
 FT Misc-difference 2486
 FT /label= Val, Met
 FT Misc-difference 2488
 FT /label= Lys, Gln
 FT Misc-difference 2844
 FT /label= Leu, Met
 FT Misc-difference 2862
 FT /label= Leu, Gln
 FT Misc-difference 2917
 FT /label= Arg, Leu
 FT Misc-difference 2968
 FT /label= Ser, Gly
 FT Misc-difference 2985
 FT /label= Cys, Arg
 FT Misc-difference 2990
 FT /label= Tyr, Cys
 XX JP0610569C-A.
 PN 19-APR-1994.
 PD 10-MAR-1992; 92JP-0051885.
 PF 10-MAR-1992; 92JP-0051885.
 XX (KAEN/) KAENNO K.
 XX WPI: 1994-153130/20.
 DR N-PSDB: AAQ63499.

FT Biocotransmissible non-A non-B hepatitis virus DNA - used for
 PT detection of hepatitis virus
 XX
 PS Claim 1: Page 8-20; 22pp: Japanese.
 XX
 CC This sequence is encoded by the genome of a blood transmissible non-A,
 CC non-B hepatitis (NANBH) virus. The cDNA sequence was isolated using the
 CC primers given in AAQ63500-35. The amplified fragments are used in the
 CC detection of hepatitis virus. The target DNA was isolated from serum
 CC of chronically infected NANBH patients who were C100 antibody-positive
 CC and HCV RNA (NS5 region) positive. Reverse transcription-PCR and PCR
 CC were performed on cDNA and the total human NANBH DNA was constructed
 CC from 23 clones.
 XX
 SQ Sequence 3010 AA;
 Query Match 97.7%; Score 43; DB 15; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 65;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 LIGCIIITSL 9
 DB 1039 LIGCIVISL 1047
 IIILIIII
 RESULT 2
 AAR50072
 ID AAR50072 standard; Protein: 273 AA.
 XX AC AAR50072;
 XX 24-MAY-1994 (first entry)
 XX NANBH virus antigenic fragment #4.
 XX Antigen; structural; non-structural; non A non B hepatitis virus;
 KW NANBH; patient; plasma; diagnosis; detection; carrier; ss.
 XX Non A Non B hepatitis virus.
 OS JP06032778-A.
 XX 15-MAR-1994.
 XX PD 01-JUN-1993; 93JP-0156087.
 XX PF 10-JUL-1992; 92JP-0207391.
 XX (KOKU-) KOKUSAI SHIYAKU KK.
 PA (SANW) SANWA KAGAKU KENKYUSHO CO.
 PA (TOFU) TONEN CORP.
 PA (TOKH-) 2H TOKYOTO RINSHO IGAKU SOGO KENKYUSHO.
 XX WPI: 1994-128677/16.
 DR N-PSDB: AAQ58817.
 XX Nucleic acid fragment coding non-A non-B hepatitis virus antigen
 PT - useful in diagnosis of NANBH patient and detection of virus
 PT carrier
 XX
 PS Claim 8; Page 18-19; 37pp: Japanese.
 XX
 CC The sequences given in AAR50068 and AAR50070-82 represent antigens of
 CC structural and non-structural regions of non A non B hepatitis virus
 CC (NANBH). The cDNA encoding these sequences were derived from the
 CC plasma of a NANBH patient by recombinant DNA techniques. These
 CC fragments are useful for the diagnosis of NANBH patients and the
 CC detection of NANBH carriers.
 XX
 SQ Sequence 273 AA;
 Query Match 95.5%; Score 42; DB 15; Length 273;
 Best Local Similarity 88.9%; Pred. No. 9.5;

```

Matches      8;  Conservative      1;  Mismatches      0;  Indels      0;  Gaps      0;

QY      1 LIGCIITSL 9
        1:|||||
Db      23 LIGCIITSL 31

RESULT 3
AAR54066
ID      AAR54066 standard; Protein: 1051 AA.
XX
AC      AAR54066;
DT      14-FEB-1995 (first entry)
DE      Non-A, non-B hepatitis virus gene #4 product.
XX
KW      Non-A, non-B hepatitis virus; NANBHV; Hepatitis C virus; HCV;
KW      core; ENV; NS1; NS2; NS3; antigen; detection.
XX
OS      Hepatitis C virus.
XX
FH      Key      Location/Qualifiers
FT      Region      1..191
FT      Region      /label= core
FT      Region      192..383
FT      Region      /label= ENV
FT      Region      384..810
FT      Region      /label= NS1
FT      Region      811..1051
FT      Region      /label= NS2-NS3
FT      Region      /note= "NS3 N-terminal"
XX
PN      JP06141870-A.
XX
XX      24-MAY-1994.
XX
PE      12-MAR-1992; 92JP-0088140.
PR      12-MAR-1992; 92JP-0088140.
XX
PA      (SANW ) SANWA KAGAKU KENKYUSHO CO.
PA      (TOFU ) TONEN CORP.
PA      (TOKR-) ZH TOKYOYO RINSHO IGAKU SOGO KENKYUSHO.
XX
XX      WPI: 1994-205828/25.
DR      N-PSDB; AAQ64068.
XX
XX      DNA coding a Non-A, non-B hepatitis virus antigen - useful for
XX      detecting HCV within serum.
XX
PS      Claim 1-5; Page 11-15; 22pp; Japanese.
XX
CC      Hepatitis C virus #4 and #6 genes were isolated (AAQ64068-69).
CC      Both genes contain the core, ENV, NS1, NS2 and NS3 regions.
CC      A core region fragment is given in AAQ64067.
XX
SQ      Sequence      1051 AA;

      Query Match      95.5%; Score 42; DB 15; Length 1051;
      Best Local Similarity      88.9%; Pred. No. 35;
      Matches      8;  Conservative      1;  Mismatches      0;  Indels      0;  Gaps      0;

QY      1 LIGCIITSL 9
        1:|||||
Db      1039 LIGCIITSL 1047

RESULT 4
AAR98361
ID      AAR98361 standard; Protein: 1051 AA.
XX
AC      AAR98361;
DT      21-JUN-1999 (first entry)
DE      Infectious hepatitis C virus genotype 1b strain HC-J4 protein.
XX
KW      HCV; infectious clone; infection; diagnosis; therapy; vaccine;
KW      screening; assay; antiviral; virucide.
XX
OS      Hepatitis C virus.
XX
PN      W05904008-A2.
XX
XX      28-JAN-1999.

```

```

XX      22-AUG-1996 (first entry)
DT
XX
DE      5'UTR/CORE/ENV/NS1/NS2/NS3 from HCV (#4).
XX
KW      Hepatitis C virus; HCV; antigen; detection; antibody.
XX
OS      Hepatitis C virus.
XX
FH      Key      Location/Qualifiers
FT      Peptide      1..191
FT      Peptide      /label= Core peptide
FT      Peptide      192..383
FT      Peptide      /label= ENV1
FT      Peptide      384..810
FT      Peptide      /label= NS1/NS2
FT      Peptide      811..1051
FT      Peptide      /label= NS2 and NS3
XX
PN      JP0713329-A.
XX
XX      23-MAY-1995.
XX
PE      18-JUN-1993; 93JP-0147944.
PR      18-JUN-1993; 93JP-0147944.
XX
PA      (TOFU ) TONEN CORP.
XX
XX      WPI: 1995-228780/29.
DR      N-PSDB; AAT30386.
XX
XX      Recombinant polypeptide comprising partial NS1 region of hepatitis
XX      non-A non-B viral antigen - used in a method for detecting
XX      antibodies against hepatitis non-A non-B virus.
XX
PS      Disclosure; Page 10-12; 15pp; Japanese.
XX
XX      The sequences given in AAR98361-62 represent the 5'UTR/CORE/ENV/NS1/NS2/
XX      NS3 protein region derived from hepatitis C virus (HCV) isolates #4
XX      and #6 respectively. The proteins partic. contain amino acids 384-
XX      495 of the HCV NS1 antigen. These protein fragments may be used in
XX      the detection of antibodies against HCV.
XX
SQ      Sequence      1051 AA;

      Query Match      95.5%; Score 42; DB 16; Length 1051;
      Best Local Similarity      88.9%; Pred. No. 35;
      Matches      8;  Conservative      1;  Mismatches      0;  Indels      0;  Gaps      0;

QY      1 LIGCIITSL 9
        1:|||||
Db      1039 LIGCIITSL 1047

RESULT 5
AAR98022
ID      AAR98022 standard; Protein: 3010 AA.
XX
AC      AAR98022;
XX
XX      21-JUN-1999 (first entry)
DE      Infectious hepatitis C virus genotype 1b strain HC-J4 protein.
XX
KW      HCV; infectious clone; infection; diagnosis; therapy; vaccine;
KW      screening; assay; antiviral; virucide.
XX
OS      Hepatitis C virus.
XX
PN      W05904008-A2.
XX
XX      28-JAN-1999.

```

XX PF 16-JUL-1998; 98WO-0514682.
 XX PR 27-JAN-1998; 98US-0014414.
 XX PR 18-JUL-1997; 97US-0033062.
 XX PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX P: Bukh J, Emerson SJ, Purcell RH, Yanagisaki M;
 XX DR WPI; 1999-132252/11.
 XX DR N-PSDB; AAX24843.
 XX PT New isolated hepatitis C virus nucleic acids - used to develop
 PT products for the diagnosis, prevention and treatment of HCV
 PT infections and for developing screening assays
 XX PS Claim 2; Fig 14G-H; 126pp; English.
 XX CC This protein is encoded by the infectious hepatitis C virus (HCV)
 CC genotype 1b strain HC-J4 genome (see AAX24833). HC-J4 was obtained
 CC from acute phase plasma of a chimpanzee infected with serum
 CC containing HC-J4/91. The infectious nucleic acid sequence can be
 CC used to produce chimeric genomes (see AAX24833) consisting of the
 CC open reading frames of infectious nucleic acid sequences of other
 CC genotypes (including genotypes 1-6) and subtypes (such as 1b, 2a,
 CC 2b, 2c, 3a, 4a-f, 5a and 6a) of HCV. The invention also relates to
 CC the introduction of mutations or deletions into infectious nucleic
 CC acid sequences in order to produce an attenuated HCV virus suitable
 CC for vaccine development. Infectious nucleic acid sequences can also
 CC be used to produce attenuated virus via passage in vitro or in vivo
 CC of the viruses produced by transfection of a host cell with the
 CC infectious nucleic acid sequence. Vaccines comprising one or more
 CC polypeptides made from the infectious nucleic acid sequence are
 CC used to immunise mammals, especially humans, against hepatitis C.
 CC The nucleic acid sequences can also be used to induce protective
 CC immunity against the virus. The nucleic acid sequences or their
 CC encoded proteases (e.g. NS3 protease) can additionally be used to
 CC develop screening assays to identify antiviral agents for HCV.
 XX SQ Sequence 3010 AA;
 Query Match 93.2%; Score 41; DB 20; Length 3010;
 Best Local Similarity 89.9%; Pred. No. 1.5e+02;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 LAGCIITSL 9
 DB 1039 VLGCIIITSL 1047
 RESULT 6
 AAB31170
 ID AAB31170 standard; Protein; 3010 AA.
 XX AC AAB31170;
 XX DT 02-APR-2001 (first entry)
 XX DE Amino acid sequence of a hepatitis C virus (HCV) clone genotype 1b.
 XX KW Chimeric virus; bovine viral diarrhoea virus; BVDV; hepatitis C virus;
 XX KW HCV; vaccine; viral inhibitor; antiviral.
 XX OS Hepatitis C virus.
 XX PN WO200075352-A2.
 XX PD 14-DEC-2000.
 XX PF 02-JUN-2000; 2000WO-US15527.
 XX PR 04-JUN-1999; 99US-0137817.

XX FA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX FI Nam J, Bukh J, Emerson SJ, Purcell RH;
 XX DR WPI; 2001-071081/08.
 XX DR N-PSDB; AAC86939.
 XX PT New nucleic acid comprising a chimeric bovine viral diarrhoea virus
 PT genome in which the (non-)structural region has been replaced by
 PT hepatitis C virus (HCV) genome useful for treating or preventing HCV
 PT signs and symptoms -
 XX PS Disclosure; Fig 4G-H; 97pp; English.
 XX CC The specification describes a nucleic acid comprising a chimeric virus
 CC genome, specifically bovine viral diarrhoea virus (BVDV) genome in which
 CC the (non-)structural region has been replaced by the (non-)structural
 CC region of a hepatitis C virus (HCV) genome. The nucleic acids comprising
 CC the chimeric virus and the chimeric virus are useful for identifying
 CC cell lines capable of supporting the replication of these chimeric
 CC viruses, in screening for neutralizing antibodies to HCV of different
 CC genotypes, in the production of HCV-BVDV virions, for the development
 CC of inactivated or attenuated vaccines to prevent HCV-BVDV in a mammal,
 CC in studying the molecular properties of HCV indirectly in vitro, and in
 CC identifying inhibitors of viral enzyme activity which would be useful
 CC as antiviral agents. Formulations or compositions comprising the
 CC chimeric virions may be used to treat or prevent the signs and symptoms
 CC of HCV. The present sequence is encoded by a HCV clone, which is used
 CC to construct chimeric nucleic acids of the invention.
 XX SQ Sequence 3010 AA;
 Query Match 93.2%; Score 41; DB 22; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 1.5e+02;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 LAGCIITSL 9
 DB 1039 VLGCIIITSL 1047
 RESULT 7
 AAB59174
 ID AAB59174 standard; Protein; 3010 AA.
 XX AC AAB59174;
 XX DT 21-MAR-2001 (first entry)
 XX DE Protein encoded by infectious Hepatitis C virus 1b genotype.
 XX KW BVDV-B; hepatitis C virus; HCV; vaccine.
 XX OS Hepatitis C virus.
 XX PN WO200075337-A1.
 XX PD 14-DEC-2000.
 XX PF 02-JUN-2000; 2000WO-US15293.
 XX PR 04-JUN-1999; 99US-0137564.
 XX PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX PI Bukh J, Yanagisaki M, Emerson SJ, Purcell RH;
 XX DR WPI; 2001-091214/10.
 XX PT New infectious nucleic acids of the GB virus-B clone, useful for
 PT indirectly studying the molecular properties of hepatitis C virus (HCV)
 PT and in developing vaccines and therapeutics for HCV -

XX PS Disclosure; Fig 7; 96pp; English.

XX CC The present invention relates to GB virus-B. The nucleic acid molecules of the invention are useful for indirectly studying the molecular properties of hepatitis C virus (HCV). The infectious nucleic acid sequence of the GB virus-B clone and the HCV/GBV-B chimeras may be used in the development of vaccines and therapeutics for HCV.

XX SQ Sequence 3010 AA:

Query Match 93.2%; Score 41; DB 22; Length 3010;
Best Local Similarity 88.5%; Pred. No. 1.5e-02;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 1 LGCIIITSL 9
:|||||

DB 1039 VLGCIIITSL 1047

RESULT 8
AA44727
ID AAY44727 standard; Protein: 184 AA.
AC AAY44727;
DI 04-MAY-2000 (first entry)
XX Hepatitis C virus NS3 catalytic domain of 1B genotype.
XX NS3 catalytic domain; NS4A peptide; NS4A-NS3 fusion construct; diagnosis;
KW serine protease; trypsin family; screening; anti-viral compound;
KW treatment; inhibitor; therapeutic.
XX Hepatitis C virus.
XX Key Location/Qualifiers
FH 1..5
FT Region
FT /note= "Residues that are deleted for the construction
FT of NS4A/NS3 fusion construct"

XX PN WO200001718-A2.
XX 13-JAN-2000.
XX 02-JUL-1999; 98WO-US15035.
XX 02-JUL-1998; 98US-0091675.
XX (UYEL) UNIV FLORIDA.
XX Dunn BM, Bukhtiyarova M;
XX WPI: 2000-182103/16.
XX Novel polypeptide comprising hepatitis C virus NS4A and NS3 domains.
XX useful for screening for compounds useful for the diagnosis and
XX treatment of hepatitis C virus.

XX PS Claim 3; Fig 1; 30pp; English.

XX CC The present protein sequence is the catalytic domain of the NS3 clone
XX from the 1B genotype, derived from hepatitis C virus. The NS3 protein
XX has a sequence relationship to members of the trypsin family of serine
XX proteases. Constructs comprising an NS4A peptide sequence covalently
XX attached to the N-terminus of NS3 catalytic domain or their fragments
XX can be used to create fusion polypeptides. This fusion polypeptide
XX contains the NS3 domain expressed in a stable, soluble form. This
XX facilitates the use of the polypeptide in direct screening of potential
XX anti-viral compounds, that are used for diagnosis and treatment of
XX hepatitis C virus infection. It is also used to screen for inhibitors of
XX serine protease activity. The polynucleotides are also useful to identify
XX diagnostic or therapeutic compounds and for recombinant production of

XX CC the fusion polypeptide.

XX SQ Sequence 184 AA:

Query Match 90.9%; Score 40; DB 21; Length 184;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 LGCIIITSL 9
:|||||

DB 15 LGCIIITSL 22

RESULT 9
AA93482
ID AAW93482 standard; Protein: 631 AA.
XX AAW93482;
XX 11-JUN-1999 (first entry)
XX HCV NS3 protein.
XX NS3; helicase domain; X-ray crystal structure; Hepatitis C virus;
KW HCV; computer program; binding pocket; three-dimensional.
XX Hepatitis C virus.
XX WO9909148-A1.
XX 25-FEB-1999.
XX 13-AUG-1998; 98WO-US16879.
XX 13-AUG-1997; 97US-0055772.
XX (VERT-) VERTEX PHARM INC.
XX Caron P, Kim J., Lin C, Morgenstern K;
XX WPI: 1999-190157/16.
XX N-PSDB; AAX21258.
XX New hepatitis C virus NS3 helicase crystals - which provide
XX molecular design techniques to identify, select and design agents
XX which bind to the helicase, particularly inhibitor compounds
XX Disclosure; Page 202-205; 224pp; English.
XX This invention relates to the X-ray crystal structure of the Hepatitis C
XX virus helicase domain. The invention describes crystallized complexes of
XX HCV helicase and an oligonucleotide. The described method is used in a
XX novel computer programme where the computer is programmed with the
XX structure coordinates of the HCV helicase oligonucleotide binding pocket
XX or the HCV helicase nucleotide triphosphate pocket wherein the computer
XX is capable of displaying a three-dimensional representation of that
XX binding pocket.

XX SQ Sequence 631 AA:

Query Match 90.9%; Score 40; DB 20; Length 631;
Best Local Similarity 100.0%; Pred. No. 49;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 LGCIIITSL 9
:|||||

DB 14 LGCIIITSL 21

RESULT 10
AA934580
ID AAR34580 standard; Protein: 3010 AA.
XX


```

AC AAR34580;
XX
XX 25-MAR-2003 (updated)
XX 25-AUG-1993 (first entry)
XX
XX Human hepatitis C virus gene encoded polyprotein.
XX
XX HCV; detection: diagnosis: vaccine; peptidc.
XX
XX Human hepatitis C virus.
XX
XX EP541089-A2.
XX
XX 12-MAY-1993.
XX
XX 05-NOV-1992; 92EP-0118974.
XX
XX 07-NOV-1991; 91JP-0312679.
XX
XX (SANW ) SANWA KAGAKU KENKYUSHO CO.
XX
XX Hayashi Y, Jomori T, Kureno M, Mitani T, Sawa K, Suzuki E;
XX
XX WPI; 1993-154074/19.
XX
XX N-PSDB; AAQ41345.
XX
XX Single or double stranded deoxyribonucleic acid for hepatitis C virus
XX
XX detection - comprises 9500 nucleotide(s) and encodes human
XX
XX hepatitis C virus gene, for diagnosis by polymerase chain reaction
XX
XX serum sample RNA deriv. and electrophoresis
XX
XX
XX Disclosure; Page 4-20; 21pp; English.
XX
XX The sequence is that encoded by a human hepatitis C virus (HCV) gene.
XX
XX A peptide comprising part of the amino acid sequence may be used as
XX
XX an antigen as part of a method of detection for human anti-HCV
XX
XX antibody or in diagnosis of human hepatitis C. It may also be used
XX
XX as an antigen in a method of prodn. of vaccine for human hepatitis C.
XX
XX (Updated on 25-MAR-2003 to correct PN field.)
XX
XX
XX Sequence 3010 AA;
XX
XX
XX Query Match 85.6%; Score 39; DB 14; Length 3010;
XX
XX Best Local Similarity 77.8%; Pred. No. 3.4e-02;
XX
XX Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
XX
XX
XX 1 LLGCIIITSJ 9
XX
XX 1039 LFGCIVTSL 1047
XX
XX
XX RESULT 11
XX
XX AAR34458
XX
XX ID AAR34458 standard; Protein; 3011 AA.
XX
XX
XX AAR34468;
XX
XX 30-JUL-1993 (first entry)
XX
XX Encoded by full-length Hepatitis C virus clone JKI 5.
XX
XX
XX HCV; non-A, non-B hepatitis virus; NANBHV; liver disease;
XX
XX polymerase chain reaction; diagnostic method.
XX
XX Hepatitis C virus.
XX
XX Key Location/Qualifiers
XX
XX Misc-difference 2414
XX
XX /note= "not defined"
XX
XX JP05068562-A.
XX
XX 23-MAR-1993.
XX
XX

```

```

XX
XX 30-MAY-1991; 91JP-0153736.
XX
XX 30-MAY-1991; 91JP-0153736.
XX
XX (SANW ) SANWA KAGAKU KENKYUSHO CO.
XX
XX WPI; 1993-130638/16.
XX
XX N-PSDB; AAQ40425.
XX
XX DNA and cDNA of hepatitis C virus - useful as probes for
XX
XX diagnosing HCV infection
XX
XX Claim 3; Page 5-18; 44pp; Japanese.
XX
XX cDNA was prepared from HCV genomic RNA. Full-length clone JKI-5
XX
XX and 14 shorter clones were isolated by PCR amplification.
XX
XX Primers/probes derived from the sequences of these clones can be used
XX
XX in diagnostic assays for HCV. See also AAQ40425-Q40439.
XX
XX
XX Sequence 3011 AA;
XX
XX
XX Query Match 88.6%; Score 39; DB 14; Length 3011;
XX
XX Best Local Similarity 77.8%; Pred. No. 3.4e-02;
XX
XX Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
XX
XX
XX 1 LLGCIIITSJ 9
XX
XX 1039 LFGCIVTSL 1047
XX
XX
XX RESULT 12
XX
XX AAR29854
XX
XX ID AAR29854 standard; Protein; 174 AA.
XX
XX
XX AAR29854;
XX
XX 25-MAR-2003 (updated)
XX
XX 26-APR-1993 (first entry)
XX
XX HCV NS2-NS4 peptide Q26-1.
XX
XX
XX Clone; polypeptide; NS2-NS4; Hepatitis C; Virus; HCV; serum; HC;
XX
XX transcriptase; cDNA; primer; allele.
XX
XX Hepatitis C virus.
XX
XX EP515313-A2.
XX
XX 16-DEC-1992.
XX
XX 11-JUN-1992; 92EP-0109812.
XX
XX 11-JUN-1991; 91JP-0139268.
XX
XX 12-JUL-1991; 91JP-0172794.
XX
XX 67-OCT-1991; 91JP-0287006.
XX
XX 16-DEC-1991; 91JP-0332329.
XX
XX 20-APR-1992; 92JP-0099957.
XX
XX (MIU ) MITSUBISHI KASEI CORP.
XX
XX Hayashi N, Honda Y, Murakami T, Seki M, Takahashi K;
XX
XX Teranishi Y;
XX
XX WPI; 1992-417213/51.
XX
XX N-PSDB; AAQ32485.
XX
XX New hepatitis C virus gene and its encoded protein - used for
XX
XX diagnosing and vaccinating against hepatitis C virus infections
XX
XX Disclosure; Page 151-52; 305pp; English.
XX
XX
XX The sequences given in AAR29852-70 are encoded by various clones which
XX

```

CC were used in the isolation of the NS2-NS4 regions of the Hepatitis C
 CC Virus (HCV) gene of the invention (see also: AAR29660, AAR29559-60 and
 CC AAR29493-51). These RNA sequences were isolated from the serum of a
 CC patient suffering from hepatitis C (HC). The isolated RNA sequences
 CC were converted into cDNA using transcriptase in the presence of one
 CC of the primer sequences given in AAG32578-79. The sequences were then
 CC amplified using primer pairs. The cDNA sequences isolated represent
 CC different alleles of the same region of the HCV gene. Sequence
 CC comparisons of these clones showed that it is possible for a patient
 CC to carry more than one HCV strain at one time. See also AAG32436.
 CC (Updated on 25-MAR-2003 to correct PN field.)

XX Sequence 174 AA:

Query Match 84.1% Score 37; DB 13; Length 174;
 Best Local Similarity 88.9%; Pred. No. 49;
 Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ILGCKITSL 9

Db 55 LLGCKITSL 64

RESULT 13

AAV17889
 ID AAV17886 standard; Protein: 215 AA.

XX AAV17889;

XX 07-SEP-1999 (first entry)

XX HCV NS4A-NS3 complex SEQ ID NO:50.

XX HCV; hepatitis C virus; single chain recombinant complex; linker;
 KW NS3 protease; NS4A cofactor; NS4A-NS3 complex; serine protease;
 KW hydrophobic domain; covalent complex; detection; inhibitor.

OS Hepatitis C virus.

OS Synthetic.

XX WO9928482-A2.

XX 10-JUN-1999.

XX 24-NOV-1998; 98WO-US24528.

XX 28-JUL-1998; 98US-0094331.

XX 28-NOV-1997; 97US-0067315.

XX (SCHE) SCHERING CORP.

PI Malcolm BA, Taremi SS, Weber PC, Yao N;

XX WPI: 1999-385385/32.

XX New hepatitis C virus covalent complexes

XX Example 1; Page 119; 21pp; English.

XX The present invention describes a covalent hepatitis C virus (HCV)
 CC NS4A-NS3 complex comprising a central hydrophobic domain of native HCV
 CC NS4A peptide, a linker, and an HCV NS3 serine protease domain, where the
 CC hydrophobic domain of native HCV NS4A peptide is tethered by the linker
 CC to the amino terminus of the HCV NS3 protease domain. The present
 CC sequence represents an example of the above complex. The covalent
 CC NS4A-NS3 complexes are useful for structural determination and
 CC determination of mode of binding of HCV inhibitors by NMR spectroscopy.
 CC They can also be used for detecting inhibitors of the protease activity,
 CC the helicase activity and the ATPase activity of NS3. The covalent
 CC NS4A-NS3 complexes are more soluble, stable and active than the non
 CC covalent protease-peptide complexes previously available.

XX Sequence 213 AA:

Query Match 84.1% Score 37; DB 20; Length 213;
 Best Local Similarity 88.9%; Pred. No. 60;
 Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ILGCKITSL 9

Db 47 LLGCKITSL 55

RESULT 14

AAV17890
 ID AAV17890 standard; Protein: 215 AA.

XX AAV17890;

XX 07-SEP-1999 (first entry)

XX HCV NS4A-NS3 complex SEQ ID NO:57.

XX HCV; hepatitis C virus; single chain recombinant complex; linker;
 KW NS3 protease; NS4A cofactor; NS4A-NS3 complex; serine protease;
 KW hydrophobic domain; covalent complex; detection; inhibitor.

OS Hepatitis C virus.

OS Synthetic.

XX WO9928482-A2.

XX 10-JUN-1999.

XX 24-NOV-1998; 98WO-US24528.

XX 28-JUL-1998; 98US-0094331.

XX 28-NOV-1997; 97US-0067315.

XX (SCHE) SCHERING CORP.

PI Malcolm BA, Taremi SS, Weber PC, Yao N;

XX WPI: 1999-385385/32.

XX New hepatitis C virus covalent complexes

XX Example 1; Page 122-123; 21pp; English.

XX The present invention describes a covalent hepatitis C virus (HCV)
 CC NS4A-NS3 complex comprising a central hydrophobic domain of native HCV
 CC NS4A peptide, a linker, and an HCV NS3 serine protease domain, where the
 CC hydrophobic domain of native HCV NS4A peptide is tethered by the linker
 CC to the amino terminus of the HCV NS3 protease domain. The present
 CC sequence represents an example of the above complex. The covalent
 CC NS4A-NS3 complexes are useful for structural determination and
 CC determination of mode of binding of HCV inhibitors by NMR spectroscopy.
 CC They can also be used for detecting inhibitors of the protease activity,
 CC the helicase activity and the ATPase activity of NS3. The covalent
 CC NS4A-NS3 complexes are more soluble, stable and active than the non-
 CC covalent protease-peptide complexes previously available.

XX Sequence 215 AA:

Query Match 84.1% Score 37; DB 20; Length 215;
 Best Local Similarity 88.9%; Pred. No. 60;
 Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ILGCKITSL 9

Db 47 LLGCKITSL 55

RESULT 15

AAV17879
 ID AAV17879 standard; Protein: 216 AA.

```

XX  RAY17879;
AC
XX  07-SEP-1999 (first entry)
XX
XX  HCV NS4A-NS3 complex SEQ ID NO:3.
XX
XX  HCV: hepatitis C virus; single chain recombinant complex; linker:
XX  NS3 protease; NS4A cofactor; NS4A-NS3 complex; serine protease;
XX  hydrophobic domain; covalent complex; detection; inhibitor.
XX
XX  Hepatitis C virus.
XX  Synthetic.
XX
XX  WO9928482-A2.
XX
XX  10-JUN-1999.
XX
XX  24-NOV-1998; 98WO-0524528.
XX
XX  28-JUL-1998; 98US-0034331.
XX  28-NOV-1997; 97US-0067315.
XX
XX  (SCHE ) SCHERING CORP.
XX
XX  Malcolm BA, Taremi SS, Weber PC, Yao N;
XX
XX  WP: 1999-385385/32.
XX
XX  New hepatitis C virus covalent complexes
XX
XX  Claim 6: Page 75-76; 21pp; English.
XX
XX  The present invention describes a covalent hepatitis C virus (HCV)
XX  NS4A-NS3 complex comprising a central hydrophobic domain of native HCV
XX  NS4A peptide, a linker, and an HCV NS3 serine protease domain, where the
XX  hydrophobic domain of native HCV NS4A peptide is tethered by the linker
XX  to the amino terminus of the HCV NS3 protease domain. The present
XX  sequence represents a specifically claimed example of the above
XX  complex. The covalent NS4A-NS3 complexes are useful for structural
XX  determination and determination of mode of binding of HCV inhibitors by
XX  NMR spectroscopy. They can also be used for detecting inhibitors of the
XX  protease activity, the helicase activity and the ATPase activity of NS3.
XX  The covalent NS4A-NS3 complexes are more soluble, stable and active than
XX  the non-covalent protease-peptide complexes previously available.
XX
SQ  Sequence 216 AA:

Query Match 84.1% Score 37; DB 20; Length 216;
Best Local Similarity 88.9%; Pred. No. 61;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY  1 ILGCLITSL 9
Db  1111111
48 LGGCKTSL 56

Search completed: September 29, 2003, 19:07:02;
Job time : 3.5 secs

```

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: September 29, 2003, 19:03:22 : Search time 13 seconds
(without alignments)

48,084 Million cell updates/sec

Title: US-09-965-594-1_COPY_14_22

Perfect score: 44

Sequence: 1 LLGCIITSL 9

Scoring table: BLOSUM62

Gapop 10.0, Gapext 0.5

Searched: 29330w seqs, 5616062 residues

Total number of hits satisfying chosen parameters: 29330

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 96%

Listing first 45 summaries

Database: PIR761*

1: pir1*

2: pir2*

3: pir3*

4: pir4*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	43	97.7	3010	1 A45573	genome polyprotein
2	40	90.9	3010	1 GNMV1W	genome polyprotein
3	39	88.6	3010	1 S18030	genome polyprotein
4	38	86.4	231	2 I16938	hypothetical prote
5	37	84.1	851	2 G71435	hypothetical prote
6	36	81.8	403	2 B69338	conserved hypotet
7	35	79.5	653	2 A49722	endoglin precursor
8	34	77.3	179	2 A01917	hypothetical prote
9	34	77.3	338	2 H86372	protein F508.40.1
10	34	77.3	345	2 G86372	protein F508.39.1
11	34	77.3	428	2 PC4163	toxin-co-regulated
12	34	77.3	946	2 S48255	probable membrane
13	34	77.3	2783	2 T31431	inositol 1,4,5-tri
14	33	75.0	350	1 S00755	pleckstrin - human
15	33	75.0	397	2 D83311	conserved hypotet
16	33	75.0	429	2 A81463	flagellar biosynth
17	33	75.0	441	2 A59744	beta-lactamase hom
18	33	75.0	472	2 G71503	probable replicati
19	33	75.0	525	2 T34544	hypothetical prote
20	33	75.0	1056	2 T02930	lysine-ketoglutarate
21	33	75.0	1089	1 PF00GA	platelet-derived g
22	32	72.7	158	2 H90459	hypothetical prote
23	32	72.7	258	2 A90459	Sec-independent pr
24	32	72.7	301	1 S12864	retinal isomerase
25	32	72.7	323	2 C89045	protein AC219.5.1
26	32	72.7	330	2 C69649	2-keto-3-deoxygluc
27	32	72.7	348	2 C81994	aldose 1-epimerase
28	32	72.7	367	2 C82943	ferri-chrome ABC tr
29	32	72.7	393	2 D65238	hypothetical_44.4

30	32	72.7	399	2 A60088	postsynaptic membr
31	32	72.7	417	2 A82984	acetyl-CoA C-acety
32	32	72.7	417	2 E98299	probable acyl-CoA
33	32	72.7	419	2 JC4522	alpha-galactosidas
34	32	72.7	438	2 AD0481	probable glucanase
35	32	72.7	442	2 D90122	putative helicase
36	32	72.7	444	2 PD0001	protein-glutamine
37	32	72.7	501	2 T45955	hypothetical prote
38	32	72.7	519	2 T43533	hexose transporter
39	32	72.7	531	2 T40480	hectose transporter
40	32	72.7	573	2 H85313	protein F2H15.1C.1
41	32	72.7	620	2 T13460	hypothetical prote
42	32	72.7	691	1 A29996	protein-glutamine
43	32	72.7	928	2 T36419	Sec76 domain prote
44	32	72.7	1064	2 T05195	saccharopine dehyd
45	32	72.7	1074	2 UC5928	semaphorin F precu

ALIGNMENTS

RESULT 1

A45573

genome polyprotein - hepatitis C virus (strain JT)

N:Contains: capsid protein C; envelope protein M; hepatitis virus (EC 3.4.21.98) (nonst protein NS4a; nonstructural protein NS4b; nonstructural protein NS5

C:Species: hepatitis C virus

C:Date: 19-May-2000 #sequence_revision: 19-May-2000 #text_change: 19-Jan-2001

C:Accession: A45573

R:Tanaka, T.; Kato, N.; Nakagawa, M.; Ootsuyama, Y.; Cho, M.J.; Nakazawa, T.; Hijik

Virus Res. 23, 39-53, 1992

A:Title: Molecular cloning of hepatitis C virus genome from a single Japanese carri

A:Reference number: A45573; MUID:92295714; PMID:1318627

A:Accession: A45573

A>Status: preliminary

A:Molecule type: DNA

A:Residues: 1-3010 <FAN>

A:Cross-references: GB:D11168; GB:D01171; NID:9221612; PIGN:BAA01943.1; PID:922161

A:Experimental source: HCV-JT

A:Note: sequence extracted from NCBI backbone (NCBIN:106206, NCBI:P:106207)

C:Superfamily: hepatitis C virus genome polyprotein

C:Keywords: ATP; glycoprotein; hydrolase; nucleotide binding; P-loop; polyprotein;

F:2 115/Product: capsid protein C #status predicted <CPC>

F:115-191/Product: envelope protein M #status predicted <EPM>

F:122-389/Product: major envelope protein E #status predicted <MEE>

F:353-729/Product: nonstructural protein NS1 #status predicted <NS1>

F:730-1006/Product: nonstructural protein NS2 #status predicted <NS2>

F:1007-1615/Product: hepatitis virus #status predicted <NS3>

F:1230-1237/Region: nucleotide-binding motif A (P-loop)

F:1312-1317/Region: nucleotide-binding motif B

F:1316-1319/Region: DEXH motif

F:1616-1862/Product: nonstructural protein NS4a #status predicted <NS4A>

F:1863-2013/Product: nonstructural protein NS5 #status predicted <NS5>

F:2014-3010/Product: nonstructural protein NS5 #status predicted <NS5>

Query Match 97.7% Score 43; DB 1; Length 3010;

Best Local Similarity 88.9%; Preq. No. 11;

Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Cy 1 LLGCIITSL 9

DB 1039 LLGCIITSL 1047

RESULT 2

GNMV1W

genome polyprotein - hepatitis C virus (strain Taiwan)

N:Contains: capsid protein C; envelope protein M; hepatitis virus (EC 3.4.21.98) (nonst protein NS4a; nonstructural protein NS4b; nonstructural protein NS5

C:Species: hepatitis C virus

A:Note: host Homo sapiens (man)

C:Date: 31-Dec-1992 #sequence_revision: 31-Dec-1992 #text_change: 19-Jan-2001

C:Accession: A40244

R:Chen, P.-J.; Lin, M.-H.; Tai, K.-F.; Liu, P.-C.; Lin, C.-J.; Chen, D.-S.
 Virology 188, 102-113, 1992
 A:Title: The Taiwanese hepatitis C virus genome: sequence determination and mapping the
 A:Reference number: A40244; MUID:92230206; PMID:1314443
 A:Accession: A40244
 A:Molecule type: genomic RNA
 A:Residues: 1-3010 <CH>
 A:Cross-references: GR:M84754
 C:Superfamily: Hepatitis C virus genome polyprotein
 C:Keywords: ATP; capsid protein; envelope protein; glycoprotein; hydrolase; nonstructural
 F:1115/Product: capsid protein C #status predicted <CP>
 F:116-191/Product: envelope protein E #status predicted <EP>
 F:192-389/Product: major envelope protein NS1 #status predicted <ME>
 F:390-729/Product: nonstructural protein NS2 #status predicted <NS2>
 F:730-1006/Product: nonstructural protein NS3 #status predicted <NS3>
 F:1007-1615/Product: nonstructural protein NS4 #status predicted <NS4>
 F:1230-1237/Region: nucleotide-binding motif A (P-loop)
 F:1230-1237/Region: nucleotide-binding motif B
 F:1312-1317/Region: DEXH motif
 F:1316-1319/Region: DEXH motif
 F:1616-1862/Product: nonstructural protein NS4a #status predicted <NS4>
 F:1863-2013/Product: nonstructural protein NS4b #status predicted <NS4b>
 F:2014-3010/Product: nonstructural protein NS5 #status predicted <NS5>
 F:196,209,233,234,250,305,325,417,423,430,448,532,540,556,576,623,645,1253,1255,2641,2673
 Query Match 90.9%; Score 40; DB 1; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 40;
 Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 LLGCIITSL 9
 DB 1039 LF8CIIITSL 1047
 RESULT 3
 S18030
 genome polyprotein - hepatitis C virus (isolate JKI)
 N:Contents: capsid protein C; envelope protein E; hepatitis C virus (isolate JKI) (nonstructural
 protein NS4a; nonstructural protein NS4b; nonstructural protein NS5
 C:Species: hepatitis C virus
 A:Variety: isolate JKI
 C:Date: 19-May-2000 #sequence_revision 19-May-2000 #text_change 23-Mar-2001
 C:Accession: S18030; S33570; A48332; S18029
 R:Honda, M.; Kaneko, S.; Masashi, U.; Kobayashi, K.; Murakami, S.
 Submitted to the EMBL Data Library, September 1999
 A:Description: A whole genome of hepatitis C virus cDNA was isolated from a single patient
 A:Reference number: S18028
 A:Accession: S18030
 A:Molecule type: genomic RNA
 A:Residues: 1-3010 <CH>
 A:Cross-references: EMBL:X51596; NID:659478; PDB:1A4A; PDB:1A4B; PDB:1A4C; PDB:1A4D; PDB:1A4E; PDB:1A4F; PDB:1A4G; PDB:1A4H; PDB:1A4I; PDB:1A4J; PDB:1A4K; PDB:1A4L; PDB:1A4M; PDB:1A4N; PDB:1A4O; PDB:1A4P; PDB:1A4Q; PDB:1A4R; PDB:1A4S; PDB:1A4T; PDB:1A4U; PDB:1A4V; PDB:1A4W; PDB:1A4X; PDB:1A4Y; PDB:1A4Z; PDB:1A50; PDB:1A51; PDB:1A52; PDB:1A53; PDB:1A54; PDB:1A55; PDB:1A56; PDB:1A57; PDB:1A58; PDB:1A59; PDB:1A60; PDB:1A61; PDB:1A62; PDB:1A63; PDB:1A64; PDB:1A65; PDB:1A66; PDB:1A67; PDB:1A68; PDB:1A69; PDB:1A70; PDB:1A71; PDB:1A72; PDB:1A73; PDB:1A74; PDB:1A75; PDB:1A76; PDB:1A77; PDB:1A78; PDB:1A79; PDB:1A80; PDB:1A81; PDB:1A82; PDB:1A83; PDB:1A84; PDB:1A85; PDB:1A86; PDB:1A87; PDB:1A88; PDB:1A89; PDB:1A90; PDB:1A91; PDB:1A92; PDB:1A93; PDB:1A94; PDB:1A95; PDB:1A96; PDB:1A97; PDB:1A98; PDB:1A99; PDB:1B00; PDB:1B01; PDB:1B02; PDB:1B03; PDB:1B04; PDB:1B05; PDB:1B06; PDB:1B07; PDB:1B08; PDB:1B09; PDB:1B10; PDB:1B11; PDB:1B12; PDB:1B13; PDB:1B14; PDB:1B15; PDB:1B16; PDB:1B17; PDB:1B18; PDB:1B19; PDB:1B20; PDB:1B21; PDB:1B22; PDB:1B23; PDB:1B24; PDB:1B25; PDB:1B26; PDB:1B27; PDB:1B28; PDB:1B29; PDB:1B30; PDB:1B31; PDB:1B32; PDB:1B33; PDB:1B34; PDB:1B35; PDB:1B36; PDB:1B37; PDB:1B38; PDB:1B39; PDB:1B40; PDB:1B41; PDB:1B42; PDB:1B43; PDB:1B44; PDB:1B45; PDB:1B46; PDB:1B47; PDB:1B48; PDB:1B49; PDB:1B50; PDB:1B51; PDB:1B52; PDB:1B53; PDB:1B54; PDB:1B55; PDB:1B56; PDB:1B57; PDB:1B58; PDB:1B59; PDB:1B60; PDB:1B61; PDB:1B62; PDB:1B63; PDB:1B64; PDB:1B65; PDB:1B66; PDB:1B67; PDB:1B68; PDB:1B69; PDB:1B70; PDB:1B71; PDB:1B72; PDB:1B73; PDB:1B74; PDB:1B75; PDB:1B76; PDB:1B77; PDB:1B78; PDB:1B79; PDB:1B80; PDB:1B81; PDB:1B82; PDB:1B83; PDB:1B84; PDB:1B85; PDB:1B86; PDB:1B87; PDB:1B88; PDB:1B89; PDB:1B90; PDB:1B91; PDB:1B92; PDB:1B93; PDB:1B94; PDB:1B95; PDB:1B96; PDB:1B97; PDB:1B98; PDB:1B99; PDB:1C00; PDB:1C01; PDB:1C02; PDB:1C03; PDB:1C04; PDB:1C05; PDB:1C06; PDB:1C07; PDB:1C08; PDB:1C09; PDB:1C10; PDB:1C11; PDB:1C12; PDB:1C13; PDB:1C14; PDB:1C15; PDB:1C16; PDB:1C17; PDB:1C18; PDB:1C19; PDB:1C20; PDB:1C21; PDB:1C22; PDB:1C23; PDB:1C24; PDB:1C25; PDB:1C26; PDB:1C27; PDB:1C28; PDB:1C29; PDB:1C30; PDB:1C31; PDB:1C32; PDB:1C33; PDB:1C34; PDB:1C35; PDB:1C36; PDB:1C37; PDB:1C38; PDB:1C39; PDB:1C40; PDB:1C41; PDB:1C42; PDB:1C43; PDB:1C44; PDB:1C45; PDB:1C46; PDB:1C47; PDB:1C48; PDB:1C49; PDB:1C50; PDB:1C51; PDB:1C52; PDB:1C53; PDB:1C54; PDB:1C55; PDB:1C56; PDB:1C57; PDB:1C58; PDB:1C59; PDB:1C60; PDB:1C61; PDB:1C62; PDB:1C63; PDB:1C64; PDB:1C65; PDB:1C66; PDB:1C67; PDB:1C68; PDB:1C69; PDB:1C70; PDB:1C71; PDB:1C72; PDB:1C73; PDB:1C74; PDB:1C75; PDB:1C76; PDB:1C77; PDB:1C78; PDB:1C79; PDB:1C80; PDB:1C81; PDB:1C82; PDB:1C83; PDB:1C84; PDB:1C85; PDB:1C86; PDB:1C87; PDB:1C88; PDB:1C89; PDB:1C90; PDB:1C91; PDB:1C92; PDB:1C93; PDB:1C94; PDB:1C95; PDB:1C96; PDB:1C97; PDB:1C98; PDB:1C99; PDB:1D00; PDB:1D01; PDB:1D02; PDB:1D03; PDB:1D04; PDB:1D05; PDB:1D06; PDB:1D07; PDB:1D08; PDB:1D09; PDB:1D10; PDB:1D11; PDB:1D12; PDB:1D13; PDB:1D14; PDB:1D15; PDB:1D16; PDB:1D17; PDB:1D18; PDB:1D19; PDB:1D20; PDB:1D21; PDB:1D22; PDB:1D23; PDB:1D24; PDB:1D25; PDB:1D26; PDB:1D27; PDB:1D28; PDB:1D29; PDB:1D30; PDB:1D31; PDB:1D32; PDB:1D33; PDB:1D34; PDB:1D35; PDB:1D36; PDB:1D37; PDB:1D38; PDB:1D39; PDB:1D40; PDB:1D41; PDB:1D42; PDB:1D43; PDB:1D44; PDB:1D45; PDB:1D46; PDB:1D47; PDB:1D48; PDB:1D49; PDB:1D50; PDB:1D51; PDB:1D52; PDB:1D53; PDB:1D54; PDB:1D55; PDB:1D56; PDB:1D57; PDB:1D58; PDB:1D59; PDB:1D60; PDB:1D61; PDB:1D62; PDB:1D63; PDB:1D64; PDB:1D65; PDB:1D66; PDB:1D67; PDB:1D68; PDB:1D69; PDB:1D70; PDB:1D71; PDB:1D72; PDB:1D73; PDB:1D74; PDB:1D75; PDB:1D76; PDB:1D77; PDB:1D78; PDB:1D79; PDB:1D80; PDB:1D81; PDB:1D82; PDB:1D83; PDB:1D84; PDB:1D85; PDB:1D86; PDB:1D87; PDB:1D88; PDB:1D89; PDB:1D90; PDB:1D91; PDB:1D92; PDB:1D93; PDB:1D94; PDB:1D95; PDB:1D96; PDB:1D97; PDB:1D98; PDB:1D99; PDB:1E00; PDB:1E01; PDB:1E02; PDB:1E03; PDB:1E04; PDB:1E05; PDB:1E06; PDB:1E07; PDB:1E08; PDB:1E09; PDB:1E10; PDB:1E11; PDB:1E12; PDB:1E13; PDB:1E14; PDB:1E15; PDB:1E16; PDB:1E17; PDB:1E18; PDB:1E19; PDB:1E20; PDB:1E21; PDB:1E22; PDB:1E23; PDB:1E24; PDB:1E25; PDB:1E26; PDB:1E27; PDB:1E28; PDB:1E29; PDB:1E30; PDB:1E31; PDB:1E32; PDB:1E33; PDB:1E34; PDB:1E35; PDB:1E36; PDB:1E37; PDB:1E38; PDB:1E39; PDB:1E40; PDB:1E41; PDB:1E42; PDB:1E43; PDB:1E44; PDB:1E45; PDB:1E46; PDB:1E47; PDB:1E48; PDB:1E49; PDB:1E50; PDB:1E51; PDB:1E52; PDB:1E53; PDB:1E54; PDB:1E55; PDB:1E56; PDB:1E57; PDB:1E58; PDB:1E59; PDB:1E60; PDB:1E61; PDB:1E62; PDB:1E63; PDB:1E64; PDB:1E65; PDB:1E66; PDB:1E67; PDB:1E68; PDB:1E69; PDB:1E70; PDB:1E71; PDB:1E72; PDB:1E73; PDB:1E74; PDB:1E75; PDB:1E76; PDB:1E77; PDB:1E78; PDB:1E79; PDB:1E80; PDB:1E81; PDB:1E82; PDB:1E83; PDB:1E84; PDB:1E85; PDB:1E86; PDB:1E87; PDB:1E88; PDB:1E89; PDB:1E90; PDB:1E91; PDB:1E92; PDB:1E93; PDB:1E94; PDB:1E95; PDB:1E96; PDB:1E97; PDB:1E98; PDB:1E99; PDB:1F00; PDB:1F01; PDB:1F02; PDB:1F03; PDB:1F04; PDB:1F05; PDB:1F06; PDB:1F07; PDB:1F08; PDB:1F09; PDB:1F10; PDB:1F11; PDB:1F12; PDB:1F13; PDB:1F14; PDB:1F15; PDB:1F16; PDB:1F17; PDB:1F18; PDB:1F19; PDB:1F20; PDB:1F21; PDB:1F22; PDB:1F23; PDB:1F24; PDB:1F25; PDB:1F26; PDB:1F27; PDB:1F28; PDB:1F29; PDB:1F30; PDB:1F31; PDB:1F32; PDB:1F33; PDB:1F34; PDB:1F35; PDB:1F36; PDB:1F37; PDB:1F38; PDB:1F39; PDB:1F40; PDB:1F41; PDB:1F42; PDB:1F43; PDB:1F44; PDB:1F45; PDB:1F46; PDB:1F47; PDB:1F48; PDB:1F49; PDB:1F50; PDB:1F51; PDB:1F52; PDB:1F53; PDB:1F54; PDB:1F55; PDB:1F56; PDB:1F57; PDB:1F58; PDB:1F59; PDB:1F60; PDB:1F61; PDB:1F62; PDB:1F63; PDB:1F64; PDB:1F65; PDB:1F66; PDB:1F67; PDB:1F68; PDB:1F69; PDB:1F70; PDB:1F71; PDB:1F72; PDB:1F73; PDB:1F74; PDB:1F75; PDB:1F76; PDB:1F77; PDB:1F78; PDB:1F79; PDB:1F80; PDB:1F81; PDB:1F82; PDB:1F83; PDB:1F84; PDB:1F85; PDB:1F86; PDB:1F87; PDB:1F88; PDB:1F89; PDB:1F90; PDB:1F91; PDB:1F92; PDB:1F93; PDB:1F94; PDB:1F95; PDB:1F96; PDB:1F97; PDB:1F98; PDB:1F99; PDB:1G00; PDB:1G01; PDB:1G02; PDB:1G03; PDB:1G04; PDB:1G05; PDB:1G06; PDB:1G07; PDB:1G08; PDB:1G09; PDB:1G10; PDB:1G11; PDB:1G12; PDB:1G13; PDB:1G14; PDB:1G15; PDB:1G16; PDB:1G17; PDB:1G18; PDB:1G19; PDB:1G20; PDB:1G21; PDB:1G22; PDB:1G23; PDB:1G24; PDB:1G25; PDB:1G26; PDB:1G27; PDB:1G28; PDB:1G29; PDB:1G30; PDB:1G31; PDB:1G32; PDB:1G33; PDB:1G34; PDB:1G35; PDB:1G36; PDB:1G37; PDB:1G38; PDB:1G39; PDB:1G40; PDB:1G41; PDB:1G42; PDB:1G43; PDB:1G44; PDB:1G45; PDB:1G46; PDB:1G47; PDB:1G48; PDB:1G49; PDB:1G50; PDB:1G51; PDB:1G52; PDB:1G53; PDB:1G54; PDB:1G55; PDB:1G56; PDB:1G57; PDB:1G58; PDB:1G59; PDB:1G60; PDB:1G61; PDB:1G62; PDB:1G63; PDB:1G64; PDB:1G65; PDB:1G66; PDB:1G67; PDB:1G68; PDB:1G69; PDB:1G70; PDB:1G71; PDB:1G72; PDB:1G73; PDB:1G74; PDB:1G75; PDB:1G76; PDB:1G77; PDB:1G78; PDB:1G79; PDB:1G80; PDB:1G81; PDB:1G82; PDB:1G83; PDB:1G84; PDB:1G85; PDB:1G86; PDB:1G87; PDB:1G88; PDB:1G89; PDB:1G90; PDB:1G91; PDB:1G92; PDB:1G93; PDB:1G94; PDB:1G95; PDB:1G96; PDB:1G97; PDB:1G98; PDB:1G99; PDB:1H00; PDB:1H01; PDB:1H02; PDB:1H03; PDB:1H04; PDB:1H05; PDB:1H06; PDB:1H07; PDB:1H08; PDB:1H09; PDB:1H10; PDB:1H11; PDB:1H12; PDB:1H13; PDB:1H14; PDB:1H15; PDB:1H16; PDB:1H17; PDB:1H18; PDB:1H19; PDB:1H20; PDB:1H21; PDB:1H22; PDB:1H23; PDB:1H24; PDB:1H25; PDB:1H26; PDB:1H27; PDB:1H28; PDB:1H29; PDB:1H30; PDB:1H31; PDB:1H32; PDB:1H33; PDB:1H34; PDB:1H35; PDB:1H36; PDB:1H37; PDB:1H38; PDB:1H39; PDB:1H40; PDB:1H41; PDB:1H42; PDB:1H43; PDB:1H44; PDB:1H45; PDB:1H46; PDB:1H47; PDB:1H48; PDB:1H49; PDB:1H50; PDB:1H51; PDB:1H52; PDB:1H53; PDB:1H54; PDB:1H55; PDB:1H56; PDB:1H57; PDB:1H58; PDB:1H59; PDB:1H60; PDB:1H61; PDB:1H62; PDB:1H63; PDB:1H64; PDB:1H65; PDB:1H66; PDB:1H67; PDB:1H68; PDB:1H69; PDB:1H70; PDB:1H71; PDB:1H72; PDB:1H73; PDB:1H74; PDB:1H75; PDB:1H76; PDB:1H77; PDB:1H78; PDB:1H79; PDB:1H80; PDB:1H81; PDB:1H82; PDB:1H83; PDB:1H84; PDB:1H85; PDB:1H86; PDB:1H87; PDB:1H88; PDB:1H89; PDB:1H90; PDB:1H91; PDB:1H92; PDB:1H93; PDB:1H94; PDB:1H95; PDB:1H96; PDB:1H97; PDB:1H98; PDB:1H99; PDB:1I00; PDB:1I01; PDB:1I02; PDB:1I03; PDB:1I04; PDB:1I05; PDB:1I06; PDB:1I07; PDB:1I08; PDB:1I09; PDB:1I10; PDB:1I11; PDB:1I12; PDB:1I13; PDB:1I14; PDB:1I15; PDB:1I16; PDB:1I17; PDB:1I18; PDB:1I19; PDB:1I20; PDB:1I21; PDB:1I22; PDB:1I23; PDB:1I24; PDB:1I25; PDB:1I26; PDB:1I27; PDB:1I28; PDB:1I29; PDB:1I30; PDB:1I31; PDB:1I32; PDB:1I33; PDB:1I34; PDB:1I35; PDB:1I36; PDB:1I37; PDB:1I38; PDB:1I39; PDB:1I40; PDB:1I41; PDB:1I42; PDB:1I43; PDB:1I44; PDB:1I45; PDB:1I46; PDB:1I47; PDB:1I48; PDB:1I49; PDB:1I50; PDB:1I51; PDB:1I52; PDB:1I53; PDB:1I54; PDB:1I55; PDB:1I56; PDB:1I57; PDB:1I58; PDB:1I59; PDB:1I60; PDB:1I61; PDB:1I62; PDB:1I63; PDB:1I64; PDB:1I65; PDB:1I66; PDB:1I67; PDB:1I68; PDB:1I69; PDB:1I70; PDB:1I71; PDB:1I72; PDB:1I73; PDB:1I74; PDB:1I75; PDB:1I76; PDB:1I77; PDB:1I78; PDB:1I79; PDB:1I80; PDB:1I81; PDB:1I82; PDB:1I83; PDB:1I84; PDB:1I85; PDB:1I86; PDB:1I87; PDB:1I88; PDB:1I89; PDB:1I90; PDB:1I91; PDB:1I92; PDB:1I93; PDB:1I94; PDB:1I95; PDB:1I96; PDB:1I97; PDB:1I98; PDB:1I99; PDB:1J00; PDB:1J01; PDB:1J02; PDB:1J03; PDB:1J04; PDB:1J05; PDB:1J06; PDB:1J07; PDB:1J08; PDB:1J09; PDB:1J10; PDB:1J11; PDB:1J12; PDB:1J13; PDB:1J14; PDB:1J15; PDB:1J16; PDB:1J17; PDB:1J18; PDB:1J19; PDB:1J20; PDB:1J21; PDB:1J22; PDB:1J23; PDB:1J24; PDB:1J25; PDB:1J26; PDB:1J27; PDB:1J28; PDB:1J29; PDB:1J30; PDB:1J31; PDB:1J32; PDB:1J33; PDB:1J34; PDB:1J35; PDB:1J36; PDB:1J37; PDB:1J38; PDB:1J39; PDB:1J40; PDB:1J41; PDB:1J42; PDB:1J43; PDB:1J44; PDB:1J45; PDB:1J46; PDB:1J47; PDB:1J48; PDB:1J49; PDB:1J50; PDB:1J51; PDB:1J52; PDB:1J53; PDB:1J54; PDB:1J55; PDB:1J56; PDB:1J57; PDB:1J58; PDB:1J59; PDB:1J60; PDB:1J61; PDB:1J62; PDB:1J63; PDB:1J64; PDB:1J65; PDB:1J66; PDB:1J67; PDB:1J68; PDB:1J69; PDB:1J70; PDB:1J71; PDB:1J72; PDB:1J73; PDB:1J74; PDB:1J75; PDB:1J76; PDB:1J77; PDB:1J78; PDB:1J79; PDB:1J80; PDB:1J81; PDB:1J82; PDB:1J83; PDB:1J84; PDB:1J85; PDB:1J86; PDB:1J87; PDB:1J88; PDB:1J89; PDB:1J90; PDB:1J91; PDB:1J92; PDB:1J93; PDB:1J94; PDB:1J95; PDB:1J96; PDB:1J97; PDB:1J98; PDB:1J99; PDB:1K00; PDB:1K01; PDB:1K02; PDB:1K03; PDB:1K04; PDB:1K05; PDB:1K06; PDB:1K07; PDB:1K08; PDB:1K09; PDB:1K10; PDB:1K11; PDB:1K12; PDB:1K13; PDB:1K14; PDB:1K15; PDB:1K16; PDB:1K17; PDB:1K18; PDB:1K19; PDB:1K20; PDB:1K21; PDB:1K22; PDB:1K23; PDB:1K24; PDB:1K25; PDB:1K26; PDB:1K27; PDB:1K28; PDB:1K29; PDB:1K30; PDB:1K31; PDB:1K32; PDB:1K33; PDB:1K34; PDB:1K35; PDB:1K36; PDB:1K37; PDB:1K38; PDB:1K39; PDB:1K40; PDB:1K41; PDB:1K42; PDB:1K43; PDB:1K44; PDB:1K45; PDB:1K46; PDB:1K47; PDB:1K48; PDB:1K49; PDB:1K50; PDB:1K51; PDB:1K52; PDB:1K53; PDB:1K54; PDB:1K55; PDB:1K56; PDB:1K57; PDB:1K58; PDB:1K59; PDB:1K60; PDB:1K61; PDB:1K62; PDB:1K63; PDB:1K64; PDB:1K65; PDB:1K66; PDB:1K67; PDB:1K68; PDB:1K69; PDB:1K70; PDB:1K71; PDB:1K72; PDB:1K73; PDB:1K74; PDB:1K75; PDB:1K76; PDB:1K77; PDB:1K78; PDB:1K79; PDB:1K80; PDB:1K81; PDB:1K82; PDB:1K83; PDB:1K84; PDB:1K85; PDB:1K86; PDB:1K87; PDB:1K88; PDB:1K89; PDB:1K90; PDB:1K91; PDB:1K92; PDB:1K93; PDB:1K94; PDB:1K95; PDB:1K96; PDB:1K97; PDB:1K98; PDB:1K99; PDB:1L00; PDB:1L01; PDB:1L02; PDB:1L03; PDB:1L04; PDB:1L05; PDB:1L06; PDB:1L07; PDB:1L08; PDB:1L09; PDB:1L10; PDB:1L11; PDB:1L12; PDB:1L13; PDB:1L14; PDB:1L15; PDB:1L16; PDB:1L17; PDB:1L18; PDB:1L19; PDB:1L20; PDB:1L21; PDB:1L22; PDB:1L23; PDB:1L24; PDB:1L25; PDB:1L26; PDB:1L27; PDB:1L28; PDB:1L29; PDB:1L30; PDB:1L31; PDB:1L32; PDB:1L33; PDB:1L34; PDB:1L35; PDB:1L36; PDB:1L37; PDB:1L38; PDB:1L39; PDB:1L40; PDB:1L41; PDB:1L42; PDB:1L43; PDB:1L44; PDB:1L45; PDB:1L46; PDB:1L47; PDB:1L48; PDB:1L49; PDB:1L50; PDB:1L51; PDB:1L52; PDB:1L53; PDB:1L54; PDB:1L55; PDB:1L56; PDB:1L57; PDB:1L58; PDB:1L59; PDB:1L60; PDB:1L61; PDB:1L62; PDB:1L63; PDB:1L64; PDB:1L65; PDB:1L66; PDB:1L67; PDB:1L68; PDB:1L69; PDB:1L70; PDB:1L71; PDB:1L72; PDB:1L73; PDB:1L74; PDB:1L75; PDB:1L76; PDB:1L77; PDB:1L78; PDB:1L79; PDB:1L80; PDB:1L81; PDB:1L82; PDB:1L83; PDB:1L84; PDB:1L85; PDB:1L86; PDB:1L87; PDB:1L88; PDB:1L89; PDB:1L90; PDB:1L91; PDB:1L92; PDB:1L93; PDB:1L94; PDB:1L95; PDB:1L96; PDB:1L97; PDB:1L98; PDB:1L99; PDB:1M00; PDB:1M01; PDB:1M02; PDB:1M03; PDB:1M04; PDB:1M05; PDB:1M06; PDB:1M07; PDB:1M08; PDB:1M09; PDB:1M10; PDB:1M11; PDB:1M12; PDB:1M13; PDB:1M14; PDB:1M15; PDB:1M16; PDB:1M17; PDB:1M18; PDB:1M19; PDB:1M20; PDB:1M21; PDB:1M22; PDB:1M23; PDB:1M24; PDB:1M25; PDB:1M26; PDB:1M27; PDB:1M28; PDB:1M29; PDB:1M30; PDB:1M31; PDB:1M32; PDB:1M33; PDB:1M34; PDB:1M35; PDB:1M36; PDB:1M37; PDB:1M38; PDB:1M39; PDB:1M40; PDB:1M41; PDB:1M42; PDB:1M43; PDB:1M44; PDB:1M45; PDB:1M46; PDB:1M47; PDB:1M48; PDB:1M49; PDB:1M50; PDB:1M51; PDB:1M52; PDB:1M53; PDB:1M54; PDB:1M55; PDB:1M56; PDB:1M57; PDB:1M58; PDB:1M59; PDB:1M60; PDB:1M61; PDB:1M62; PDB:1M63; PDB:1M64; PDB:1M65; PDB:1M66; PDB:1M67; PDB:1M68; PDB:1M69; PDB:1M70; PDB:1M71; PDB:1M72; PDB:1M73; PDB:1M74; PDB:1M75; PDB:1M76; PDB:1M77; PDB:1M78; PDB:1M79; PDB:1M80; PDB:1M81; PDB:1M82; PDB:1M83; PDB:1M84; PDB:1M85; PDB:1M86; PDB:1M87; PDB:1M88; PDB:1M89; PDB:1M90; PDB:1M91; PDB:1M92; PDB:1M93; PDB:1M94; PDB:1M95; PDB:1M96; PDB:1M97; PDB:1M98; PDB:1M99; PDB:1N00; PDB:1N01; PDB:1N02; PDB:1N03; PDB:1N04; PDB:1N05; PDB:1N06; PDB:1N07; PDB:1N08; PDB:1N09; PDB:1N10; PDB:1N11; PDB:1N12; PDB:1N13; PDB:1N14; PDB:1N15; PDB:1N16; PDB:1N17; PDB:1N18; PDB:1N19; PDB:1N20; PDB:1N21; PDB:1N22; PDB:1N23; PDB:1N24; PDB:1N25; PDB:1N26; PDB:1N27; PDB:1N28; PDB:1N29; PDB:1N30; PDB:1N31; PDB:1N32; PDB:1N33; PDB:1N34; PDB:1N35; PDB:1N36; PDB:1N37; PDB:1N38; PDB:1N39; PDB:1N40; PDB:1N41; PDB:1N42; PDB:1N43; PDB:1N44; PDB:1N45; PDB:1N46; PDB:1N47; PDB:1N48; PDB:1N49; PDB:1N50; PDB:1N51; PDB:1N52; PDB:1N53; PDB:1N54; PDB:1N55; PDB:1N56; PDB:1N57; PDB:1N58; PDB:1N59; PDB:1N60; PDB:1N61; PDB:1N62; PDB:1N63; PDB:1N64; PDB:1N65; PDB:1N66; PDB:1N67; PDB:1N68; PDB:1N69; PDB:1N70; PDB:1N71; PDB:1N72; PDB:1N73; PDB:1N74; PDB:1N75; PDB:1N76; PDB:1N77; PDB:1N78; PDB:1N79; PDB:1N80; PDB:1N81; PDB:1N82; PDB:1N83; PDB:1N84; PDB:1N85; PDB:1N86; PDB:1N87; PDB:1N88; PDB:1N89; PDB:1N90; PDB:1N91; PDB:1N92; PDB:1N93; PDB:1N94; PDB:1N95; PDB:1N96; PDB:1N97; PDB:1N98; PDB:1N99; PDB:1O00; PDB:1O01; PDB:1O02; PDB:1O03; PDB:1O

Db 291 LGCTITSL 293
|||||
RESULT 6
B69338
conserved hypothetical protein AFC706 - Archaeoglobus fulgidus
C:Species: Archaeoglobus fulgidus
C>Date: 05-Dec-1997 #sequence_revision 05-Dec-1997 #text_change 22-Oct-1999
C:Accession: B69338
R:Kienk, H.P.; Clayton, R.A.; Tomb, C.F.; White, C.; Nelson, K.E.; Ketchum, K.A.; Dodson
R: Fleischmann, R.D.; Quackenbush, J.; Lee, N.H.; Sutton, G.G.; Gill, S.; Kirkness, E.F.
G:Glocke, A.; Zhou, L.; Overbeek, R.; Gocayne, J.D.; Weidman, J.F.; McDonald, L.
Nature 390, 364-370, 1997
A:Authors: Overbeek, R.; Cotton, M.D.; Springs, T.; Attlich, P.; Kaine, B.P.; Sykes, S.
Smith, H.O.; Woese, C.R.; Venter, J.C.
A:Title: The complete genome sequence of the hyperthermophilic, sulfate-reducing archae
A:Reference number: A69250; MUID:94249343; PMID:9389475
A:Accession: B69338
A>Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-403 <KL>
A:Cross-references: GB:AE001056; GR:AE000722; NID:94569379; PIDN:AA800955; PID:0254991
Query Match 81.8% Score 34; DB 2; Length 403;
Best Local Similarity 66.7% Pred. No. 41;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
QY 1 LGCTITSL 9
| | | | |
Db 330 IFGCIVTAL 338
| | | | |
RESULT 7
A49722
endoglin precursor - pig
C:Species: Sus scrofa domestica (domestic pig)
C>Date: 03-May-1994 #sequence_revision 03-May-1994 #text_change 05-Nov-1999
C:Accession: A49722; S40180
R:Yamashita, H.; Ichijo, H.; Grimsby, S.; Moran, A.; ten Dijke, P.; Miyazawa, K.
J. Biol. Chem. 269, 1995-2003, 1994
A:Title: Endoglin forms a heteromeric complex with the signaling receptors for transform
A:Reference number: A49722; MUID:94124550; PMID:8294451
A:Accession: A49722
A>Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-853 <YAM>
A:Cross-references: GR:722142; NID:9437919; PIDN:CA866273; PID:0437920
C:Keywords: homodimer; phosphoprotein; signal transduction; transmembrane protein
Query Match 79.5% Score 35; DB 2; Length 653;
Best Local Similarity 87.5% Pred. No. 54;
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 2 LGCTITSL 9
| | | | |
Db 379 LGCTITSL 385
| | | | |
RESULT 8
AD1917
hypothetical protein all0887 [imported] - Nostoc sp. (strain PCC 7120)
C:Species: Nostoc sp. PCC 7120
A:Note: Nostoc sp. strain PCC 7120 is a synonym of Anabaena sp. strain PCC 7120
C>Date: 14-Dec-2001 #sequence_revision 14-Dec-2001 #text_change 09-Dec-2002
C:Accession: AD1917
R:Kaneko, T.; Nakamura, Y.; Wolk, C.P.; Kuriu, T.; Sasamoto, S.; Watanabe, A.; Itiguchi
Nakazaki, N.; Shimpo, S.; Sugimoto, M.; Takazawa, Y.; Yamada, M.; Tabata, S.
DNA Res. 8, 205-213, 2001
A:Title: Complete Genomic Sequence of the Filamentous Nitrogen-fixing Cyanobacterium Ana
A:Reference number: AB-807; MUID:21595285; PMID:11759640
A:Accession: AD1917
A>Status: preliminary

A:Molecule type: DNA
A:Residues: 1-179 <KUR>
A:Cross-references: GB:BA000019; PIDN:BAB72844.1; PID:sl7130232; GSPDB:GN00179
A:Experimental source: strain PCC 7120
C:Genetics:
A:Gene: all0887
Query Match 77.3% Score 34; DB 2; Length 179;
Best Local Similarity 85.7% Pred. No. 49;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 LGCTITSL 7
| | | | |
Db 48 LGCTITSL 54
| | | | |
RESULT 9
H86372
protein F508.40 [imported] - Arabidopsis thaliana
C:Species: Arabidopsis thaliana (mouse-ear cress)
C>Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 31-Mar-2001
C:Accession: H86372
R:Theologis, A.; Ecker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; White, C.; A
ansen, N.F.; Hughes, B.; Huizar, L.
Nature 408, 816-820, 2000
A:Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.; K
C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, S.X.; Liu, Z.A.; Luros, J.S.; Maiti, R.; Mar
Rizzo, M.; Rooney, T.; Rowley, D.; Sakano, H.
A:Authors: Salzberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Ta
ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.
A:Title: Sequence and analysis of chromosome 1 of the plant Arabidopsis.
A:Reference number: A86141; MUID:21016719; PMID:11130712
A:Accession: H86372
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-338 <STC>
A:Cross-references: GB:AA005172; NID:94056465; PIDN:AA098039.1; GSPDB:GN00141
C:Genetics:
A:Gene: F508.40
A:Map position: 1
Query Match 77.3% Score 34; DB 2; Length 338;
Best Local Similarity 85.7% Pred. No. 83;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 3 GCLITSL 9
| | | | |
Db 136 GCLITSL 142
| | | | |
RESULT 10
S86372
protein F508.39 [imported] - Arabidopsis thaliana
C:Species: Arabidopsis thaliana (mouse-ear cress)
C>Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 31-Mar-2001
C:Accession: S86372
R:Theologis, A.; Ecker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; White, C.; A
Chin, C.W.; Chung, M.K.; Conn, L.; Conway, A.B.; Huizar, L.
Nature 408, 816-820, 2000
A:Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.; K
C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, S.X.; Liu, Z.A.; Luros, J.S.; Maiti, R.; Mar
Rizzo, M.; Rooney, T.; Rowley, D.; Sakano, H.
A:Authors: Salzberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Ta
ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.
A:Title: Sequence and analysis of chromosome 1 of the plant Arabidopsis.
A:Reference number: A86141; MUID:21016719; PMID:11130712
A:Accession: S86372
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-345 <STO>
A:Cross-references: GB:AE005172; NID:94056466; PIDN:AA098039.1; GSPDB:GN00141

C:Genetics:

A:Gene: F508.35

A:Map position: 1

Query Match 77.3%; Score 34; DB 2; Length 345;

Best Local Similarity 85.7%; Pred. No. 85;

Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 3 LGCIITS 9

II:II:II

Db 146 GCIITSL 152

RESULT 11

PC4163

toxin-co-regulated protein chain 1 - Vibrio cholerae (fragment)

C:Species: Vibrio cholerae

C:Date: 10-May-1996 #sequence_revision 19-Jul-1996 #text_change 17-Nov-2000

C:Accession: PC4163; S23261

R:Oglerman, M.A.; Voss, E.; Meaney, C.; Faust, R.; Attridge, S.R.; Manning, P.A.

Gene 170, 9-16, 1996

A:Title: Comparison of the promoter proximal regions of the toxin-co-regulated top gene

A:Reference number: JC4719; MUID:96200845; PMID:8621056

A:Accession: PC4163

A:Molecule type: DNA

A:Residues: 1-428 <OGI>

A:Cross-references: EMBL:X64098

R:Manning, P.A.

submitted to the EMBL Data Library, January 1992

A:Reference number: S23261

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-401, /RRPKN/ <MAN>

A:Cross-references: EMBL:X64098; NID:q48404; PIDN:CAA45452.1; EID:q48405

C:Comment: This is a methyl-accepting chemotaxis protein.

C:Genetics:

A:Gene: top1

Query Match

Best Local Similarity 87.5%; Score 34; DB 2; Length 428;

Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 LGCIITS 8

II:II:II

Db 11 LGCIITS 18

RESULT 12

S48255

probable membrane protein YBR086c - yeast (Saccharomyces cerevisiae)

N:Alternate names: hypothetical protein YBR086c

C:Species: Saccharomyces cerevisiae

C:Date: 03-Aug-1995 #sequence_revision 11-Aug-1995 #text_change 19-Apr-2002

C:Accession: S48255; S45954; S44670

R:Mannhaupt, G.; Stocker, R.; Ehle, S.; Vetter, I.; Feldmann, H.

Yeast 10, 1363-1381, 1994

A:Title: Analysis of a 70 kb region on the right arm of yeast chromosome II.

A:Reference number: S48255; MUID:95208357; PMID:7403426

A:Accession: S48255

A:Status: nucleic acid sequence not shown

A:Molecule type: DNA

A:Residues: 1-946 <MAN>

A:Cross-references: EMBL:X78993; NID:q475045; PIDN:CAA55593.1; PID:q476042

R:Feldmann, H.; Mannhaupt, G.; Schwarzllose, C.; Vetter, I.

submitted to the Protein Sequence Database, August 1994

A:Reference number: S45927

A:Accession: S45954

A:Molecule type: DNA

A:Residues: 1-946 <FF2>

A:Cross-references: EMBL:Z35955; NID:q536351; PID:q536352; MIPS:YBR086c

C:Genetics:

A:Gene: SGD:IST2

A:Cross-references: SGD:S0000290

A:Map position: 2R

C:Superfamily: Saccharomyces cerevisiae probable membrane protein YBR086c

C:Keywords: transmembrane protein

F:131-147/Domain: transmembrane #status predicted <TM1>

F:158-174/Domain: transmembrane #status predicted <TM2>

F:207-243/Domain: transmembrane #status predicted <TM3>

F:248-274/Domain: transmembrane #status predicted <TM4>

F:302-324/Domain: transmembrane #status predicted <TM5>

F:450-477/Domain: transmembrane #status predicted <TM6>

F:506-532/Domain: transmembrane #status predicted <TM7>

F:563-588/Domain: transmembrane #status predicted <TM8>

Query Match

Best Local Similarity 77.3%; Score 34; DB 2; Length 946;

Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 LGCIITS 8

II:II:II

Db 520 LGCIITS 527

RESULT 13

T31431

inositol 1,4,5-trisphosphate receptor, localized in plasma membrane - Panulirus arg

C:Species: Panulirus argus

C:Date: 19-May-2000 #sequence_revision 19-May-2000 #text_change 19-May-2000

C:Accession: T31431

R:Munger, S.D.; Ache, B.W.; Greenberg, R.M.

submitted to the EMBL Data Library, March 1998

A:Description: Plasma membrane localization of an olfactory inositol 1,4,5-trisphos

A:Reference number: Z21030

A:Accession: T31431

A:Status: preliminary

A:Molecule type: mRNA

A:Residues: 1-2783 <MUN>

A:Cross-references: EMBL:AF055079; NID:q3660666; PID:q3660667; PIDN:AAC61691.1

C:Superfamily: inositol-trisphosphate receptor

Query Match

Best Local Similarity 77.3%; Score 34; DB 2; Length 2783;

Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 LGCIITS 9

II:II:II

Db 2554 LLMCIIVSL 2562

RESULT 14

S00755

pleckstrin - human

N:Alternate names: p47; platelet/leukocyte C kinase substrate (pleckstrin)

C:Species: Homo sapiens (man)

C:Date: 31-Dec-1988 #sequence_revision 31-Dec-1988 #text_change 20-Apr-2000

C:Accession: S00755; A45762

R:Tyers, M.; Rachubinski, R.A.; Stewart, M.J.; Varrichio, A.M.; Short, R.G.D.; Hasl,

Nature 333, 470-473, 1988

A:Title: Molecular cloning and expression of the major protein kinase C substrate o

A:Reference number: S00755; MUID:88232910; PMID:2897630

A:Accession: S00755

A:Molecule type: mRNA

A:Residues: 1-350 <TYE>

A:Cross-references: EMBL:X07743; NID:q35517; PIDN:CAA30564.1; PID:q35518

R:Tyers, M.; Haslam, R.J.; Rachubinski, R.A.; Hatley, C.B.

J. Cell. Biochem. 40, 133-145, 1989

A:Title: Molecular analysis of pleckstrin: the major protein kinase C substrate of i

A:Reference number: A45762; MUID:89359547; PMID:2768345

A:Accession: A45762

A:Molecule type: mRNA

A:Residues: 1-350 <TY2>

A:Cross-references: GB:X07743; NID:q35517; PIDN:CAA30564.1; PID:q35518

A:Note: 92-Arg was also found

C:Genetics:

A:Gene: GDB:PIEK; p47
 A:Cross-references: GDB:383661s; OMIM:173570
 A:Map position: 2pter-2qter
 C:Superfamily: pleckstrin; pleckstrin repeat homology
 C:Keywords: phosphoprotein; signal transduction
 F:3-99/Domain: pleckstrin repeat homology <PLK1>
 F:243-345/Domain: pleckstrin repeat homology <PLK2>

Query Match 75.04; Score 33; DB 1; Length 350;
 Best Local Similarity 55.6%; Pred. No. 1.3e+02;
 Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0

QY 1 LAGCIITSL 9
 L I L L L L L
 DB 292 LRGVVTNV 305

RESULT 15
 D83311
 conserved hypothetical protein FA2562 [imported] - Pseudomonas aeruginosa (strain PA01)
 C:Species: Pseudomonas aeruginosa
 C>Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 31-Dec-2000
 C:Accession: D83311
 R:Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warren, P.; Hickey, M.J.; Br
 adman, S.; Yuan, Y.; Brody, L.L.; Collier, S.N.; Folger, K.R.; Kas, A.; Larbig, K.; Lim
 ;; Lory, S.; Olson, M.V.
 Nature 406, 959-964, 2000
 A:Title: Complete genome sequence of Pseudomonas aeruginosa PA01, an opportunistic patho
 A:Reference number: A82950; M31D:2043737; PMID:1094012
 A:Accession: D83311
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-397 <STO>
 A:Cross-references: GH:A82004895; GH:A82044831; NID:q944870; PION:AAG 6659.1; GSP:HGNC:0
 A:Experimental source: strain PA01
 C:Genetics:
 A:Gene: PA2562

Query Match 75.04; Score 33; DB 2; Length 397;
 Best Local Similarity 55.6%; Pred. No. 1.5e+02;
 Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0

QY 1 LAGCIITSL 9
 L I L L L L L
 DB 221 LRGCVLVAL 229

Search completed: September 29, 2003, 19:05:56
 Job time : 22 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: September 29, 2003, 18:58:47 ; Search time 10 Seconds
(without alignments)
42.324 Million cell updates/sec

Title: US-09-965-594-1_COPY_14_22
Perfect score: 44
Sequence: 1 LSCQIIHS 9

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 4726735 residues

Total number of hits satisfying chosen parameters: 127859

Minimum DB seq length: 0
Maximum DB seq length: 200000000
Post-processing: Minimum Match 0%
Maximum Match 99%
Listing first 45 summaries

Database : Swissprot_41*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query %	Match	Length	DB	ID	Description
1	43	97.7	310	1	POLG_HCVJT	1	P29549 h genome po
2	40	90.9	310	1	POLG_HCVJT	1	P29549 h genome po
3	36	81.8	300	1	T289_MOUSE	1	Q91414 mus muscu
4	35	79.5	653	1	EGN_PIG	1	P31776 sus scrofa
5	34	77.3	946	1	YB76_YEAST	1	P38250 saccharomy
6	33	75.0	318	1	T287_HUMAN	1	Q91414 mus muscu
7	33	75.0	350	1	PLEK_HUMAN	1	P38250 saccharomy
8	33	75.0	350	1	PLEK_MOUSE	1	Q91414 mus muscu
9	33	75.0	406	1	ARGD_PSEPK	1	Q91414 mus muscu
10	33	75.0	437	1	FLHF_PSEPK	1	P53119 pseudomonas
11	33	75.0	441	1	YB8E_BACSU	1	Q52256 pseudomonas
12	33	75.0	795	1	SYFB_SHEON	1	Q91414 mus muscu
13	33	75.0	1089	1	PGUS_HUMAN	1	Q86199 shewanella
14	33	75.0	1906	1	DICE_MOUSE	1	P10234 homo sapien
15	33	75.0	1912	1	DICE_HUMAN	1	Q84118 mus muscu
16	32	72.7	222	1	PSD9_RAT	1	Q90473 hcrpo sapien
17	32	72.7	301	1	RECS_TODPA	1	Q94755 rattus norv
18	32	72.7	330	1	KUGT_BACSU	1	P23820 todarodes p
19	32	72.7	398	1	YJGN_ECOLI	1	P58847 bacillus su
20	32	72.7	411	1	RAPS_CRICK	1	P39338 escherichia
21	32	72.7	419	1	AGAL_MOUSE	1	Q42193 gallus gall
22	32	72.7	531	1	GM2_SCHPO	1	P51569 mus muscu
23	32	72.7	689	1	TGM2_CAVCU	1	Q74969 schizosacch
24	32	72.7	928	1	YDGI_SCHPO	1	P08587 cavia cutle
25	32	72.7	1074	1	SM5A_MOUSE	1	Q10491 schizosacch
26	32	72.7	1077	1	SM5A_MOUSE	1	Q13591 homo sapien
27	31	70.5	148	1	SSRP_NEIMA	1	Q62217 mus muscu
28	31	70.5	162	1	ILJ5_BOVIN	1	Q51111 neisseria m
29	31	70.5	180	1	SP22_MOUSE	1	Q28026 bos tauris
30	31	70.5	237	1	LAPT_PASHA	1	Q9365 mus muscu
31	31	70.5	312	1	MCAL_HUMAN	1	Q12904 pasteurella
32	31	70.5	312	1	T285_MOUSE	1	Q12904 homo sapien
33	31	70.5	312	1	YM17_YEAST	1	P59530 mus muscu
							Q04516 saccharomyc

34	31	70.5	334	1	YC52_PSEAE	1	Q91492 pseudomonas
35	31	70.5	398	1	YJGN_ECO57	1	P58219 escherichia
36	31	70.5	411	1	RAPS_HUMAN	1	Q13702 homo sapien
37	31	70.5	411	1	RAPS_MOUSE	1	P12672 mus muscu
38	31	70.5	411	1	RAPS_TORCA	1	P09108 torpedo cal
39	31	70.5	417	1	PCL_RHOCA	1	C69140 rhodobacter
40	31	70.5	427	1	ETLR_PIG	1	Q29010 sus scrofa
41	31	70.5	436	1	Y326_METJA	1	O57772 methanococ
42	31	70.5	439	1	IOLF_BACSU	1	P42417 bacillus su
43	31	70.5	495	1	POLG_DEN23	1	P14339 dengue viru
44	31	70.5	556	1	PT1_STRCO	1	Q9K21 streptomyce
45	31	70.5	609	1	THI3_YEAST	1	Q07471 saccharomyc

ALIGNMENTS

RESULT 1
POLG_HCVJT
ID POLG_HCVJT STANDARD: PRT: 301C AA.
AC Q00263;
BT 01-APR-1993 (Rel. 25, Last sequence update)
DT 01-APR-1993 (Rel. 25, Last annotation update)
DE Genome polyprotein [Contains: Capsid protein C (Core protein) (P22); Envelope glycoprotein E1 (GP32) (GP35); Envelope glycoprotein E2 (GP68) (GP70) (NS1); Protein P7; Nonstructural protein NS2 (P21) (EC 3.4.22.-); Protease/helicase NS3 (P70) (Hepacivirus) (EC 3.4.21.98); Nonstructural protein NS4A (P4); Nonstructural protein NS4B (P66) (P70) (RNA-directed RNA polymerase) (EC 2.7.7.48)].
DE Hepatitis C virus (isolate HC-JT) (HCV).
OS Viruses; ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
OC Hepacivirus.
CX NCBI_TaxID=31642;
RN 111
RF SEQUENCE FROM N.A.
RX MEDLINE=92295714; Pubmed=1318527;
RA Tanaka T., Kato N., Nakagawa M., Ootsuyama Y., Cho M.J., Nakazawa T., Hilkata M., Ishimura Y., Shimotohno K.;
RT "Molecular cloning of hepatitis C virus genome from a single Japanese carrier: sequence variation within the same individual and among infected individuals."
RI Virus Res. 23:39-53(1992)
CC 1- FUNCTION: THE SMALL PROTEINS NS2A, NS2B, NS4A AND NS4B ARE HYDROPHOBIC, SUGGESTING A POSSIBLE MEMBRANE-RELATED FUNCTION. NS3 AND NS5 MAY PLAY A ROLE IN THE VIRAL RNA REPLICATION. 2- CATALYTIC ACTIVITY: Hydrolysis of four peptide bonds in the viral precursor polyprotein, commonly with Asp or Glu in the P6 position, Cys or Thr in P1 and Ser or Ala in P1'. 3- CATALYTIC ACTIVITY: N nucleoside triphosphate = N diphosphate + {RNA}(N).
CC 4- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS: PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF PROTEIN C AND MRNA.
CC 5- SIMILARITY: THE PROTEASE BELONGS TO PEPTIDASE FAMILY S29.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL Outstation - European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See http://www.isb-sib.ch/announcement/ or send an email to license@isb-sib.ch).
CC -----
CC EMBL: D11168; BAA01943.1; ..
CC PIR: A45573; A45573
CC PDB: 1A1Q; 25-MAR-98.
CC PDB: 1JXP; 14-JAN-98.
CC MEROPS: S29.001; ..
CC MEROPS: U39.001; ..
CC InterPro: IPR001410; DEAD.

DR	Pfam: PF01560; HCV_NSI; 1.	US	Mus musculus (Mouse).
DR	Pfam: PF01538; HCV_NS2; 1.	OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
DR	Pfam: PF02907; HCV_NS3; 1.	OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
DR	Pfam: PF01006; HCV_NS4a; 1.	OX	NCBI_taxonomy-10090;
DR	Pfam: PF01001; HCV_NS4b; 1.	RN	[1]
DR	Pfam: PF01506; HCV_NS5a; 1.	RP	SEQUENCE FROM N.A., AND TOPOLOGY.
DR	Pfam: PF00271; Helicase_C; 1.	RC	STRAIN-129/Svj;
DR	Pfam: PF00998; Viral_RdRP; 1.	RX	MEDLINE-20222571; PubMed-10761934;
DR	ProDom: PD185062; HCV_NSI; 1.	RA	Adler E., Hoon M.A., Mueller K.L., Chandrashekar J., Ryba N.J.P.,
DR	SMART: SM00487; DEXDC; 1.	RA	Zuker C.S.;
KW	Polypeptide; Glycoprotein; Transferase; RNA-directed RNA polymerase;	RI	"A novel family of mammalian taste receptors.";
KW	Core protein; Coat protein; Envelope protein; Helicase; ATP-binding;	RL	Cell 100:693-702(2000).
KW	Transmembrane; Nonstructural protein; Hydrolase; Serine protease;	RN	[2]
KW	3D-structure.	RP	CHARACTERIZATION.
FI	INIT_MEI 1	RX	MEDLINE-20222572; PubMed-10761935;
FI	CHAIN 1	RA	Chandrashekar J., Mueller K.L., Hoon M.A., Adler E., Peng L., Guo W.,
FI	CHAIN 116	RA	Zuker C.S., Ryba N.J.;
FI	CHAIN 191	RI	"T2Rs function as bitter taste receptors.";
FI	CHAIN 192	RL	Cell 100:703-711(2000).
FI	CHAIN 384	RN	[3]
FI	CHAIN 729	RP	REVIEW.
FI	CHAIN 730	RX	MEDLINE-22135574; PubMed-12139982;
FI	CHAIN 1007	RA	Montmayeur J.-P., Matsunami H.;
FI	CHAIN 1615	RI	"Receptors for bitter and sweet taste.";
FI	CHAIN 1862	RL	Curr. Opin. Neurobiol. 12:366-371(2002).
FI	CHAIN 1863	RN	[4]
FI	CHAIN 2014	RP	REVIEW.
FI	CHAIN 2015	RX	MEDLINE-21634924; PubMed-11696554;
FI	CHAIN 2016	RA	Margolskee R.F.;
FI	CHAIN 2017	RI	"Molecular mechanisms of bitter and sweet taste transduction.";
FI	CHAIN 2018	RL	J. Biol. Chem. 277:1-4(2002).
FI	CHAIN 2019	RN	[5]
FI	CHAIN 2020	RP	REVIEW.
FI	CHAIN 2021	RX	MEDLINE-22469025; PubMed-12581520;
FI	CHAIN 2022	RA	Zhang Y., Hoon M.A., Chandrashekar J., Mueller K.L., Cook B., Wu D.,
FI	CHAIN 2023	RI	Zuker C.S., Ryba N.J.;
FI	CHAIN 2024	RL	"Coding of sweet, bitter, and umami tastes: different receptor cells
FI	CHAIN 2025	RN	sharing similar signaling pathways.";
FI	CHAIN 2026	RP	Cell 112:293-301(2003).
FI	CHAIN 2027	CC	FUNCTION: Receptor that may play a role in the perception of
FI	CHAIN 2028	CC	bitterness and is gustducin-linked. May play a role in sensing the
FI	CHAIN 2029	CC	chemical composition of the gastrointestinal content. The activity
FI	CHAIN 2030	CC	of this receptor may stimulate alpha gustducin, mediate PLC-beta-2
FI	CHAIN 2031	CC	activation and lead to the gating of TRPM5. Functions as a ligand
FI	CHAIN 2032	CC	for cyclohexamide.
FI	CHAIN 2033	CC	CELLULAR LOCATION: Integral membrane protein.
FI	CHAIN 2034	CC	TISSUE SPECIFICITY: Expressed in subsets of taste receptor cells
FI	CHAIN 2035	CC	of the tongue and palate epithelium and exclusively in gustducin-
FI	CHAIN 2036	CC	positive cells. Expressed in gastric and duodenal tissues.
FI	CHAIN 2037	CC	MISCELLANEOUS: Most taste cells may be activated by a limited
FI	CHAIN 2038	CC	number of bitter compounds; individual taste cells can
FI	CHAIN 2039	CC	discriminate among bitter stimuli.
FI	CHAIN 2040	CC	SIMILARITY: Belongs to family T2R of G-protein coupled receptors.
FI	CHAIN 2041	CC	CAUTION: This protein was previously referred to T2R5, but due to
FI	CHAIN 2042	CC	its high similarity towards both the human and rat T2R9 sequences
FI	CHAIN 2043	CC	it is considered to be T2R9.
FI	CHAIN 2044	CC	-----
FI	CHAIN 2045	CC	This SWISS-PROT entry is copyright. It is produced through a collaboration
FI	CHAIN 2046	CC	between the Swiss Institute of Bioinformatics and the EMBL outstation
FI	CHAIN 2047	CC	the European Bioinformatics Institute. There are no restrictions on its
FI	CHAIN 2048	CC	use by non-profit institutions as long as its content is in no way
FI	CHAIN 2049	CC	modified and this statement is not removed. Usage by and for commercial
FI	CHAIN 2050	CC	entities requires a license agreement (See http://www.isb-sib.ch/announce/
FI	CHAIN 2051	CC	or send an email to license@isb-sib.ch).
FI	CHAIN 2052	CC	-----
FI	CHAIN 2053	CC	EMBL: AF27147; AAF43920.1;
FI	CHAIN 2054	CC	Pfam: PF05296; TAS2R; 1.
FI	CHAIN 2055	KW	Receptor; G-protein coupled receptor; Transmembrane.
FI	CHAIN 2056	FT	DOMAIN 1 7 EXTRACELLULAR (POTENTIAL).
FI	CHAIN 2057	FT	TRANSMEM 8 28 1 (POTENTIAL).
FI	CHAIN 2058	FT	DOMAIN 29 43 CYTOPLASMIC (POTENTIAL).
FI	CHAIN 2059	FT	TRANSMEM 44 64 2 (POTENTIAL).
FI	CHAIN 2060	FT	DOMAIN 65 87 EXTRACELLULAR (POTENTIAL).

Query Match 90.9%; Score 40; DB 1; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 21;
 Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 LLGCIITSL 9
 DB 1039 LFGCIITSL 1047

RESULT 3
 T2R9_MOUSE STANDARD; PRT; 300 AA.
 AC Q9JKT4;
 DT 15-SEP-2003 (Rel. 42, Created)
 DT 15-SEP-2003 (Rel. 42, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Taste receptor type 2 member 9 (T2R9) (Taste receptor type 2 member
 DE 5) (T2R5).
 GN TAS2R5.

```

FT TRANSMEM 88 108 3 (POTENTIAL).
FT DOMAIN 109 128 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 129 149 EXTRACELLULAR (POTENTIAL).
FT DOMAIN 150 181 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 182 202 TRANSMEM 5 (POTENTIAL).
FT DOMAIN 203 233 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 234 254 TRANSMEM 6 (POTENTIAL).
FT DOMAIN 255 259 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 260 280 TRANSMEM 7 (POTENTIAL).
FT DOMAIN 281 303 CYTOPLASMIC (POTENTIAL).
FT CARBOHYD 161 161 N-LINKED (GLCNAC...) (POTENTIAL).
SQ SEQUENCE 300 AA: 34416 MW: 92AD449C66FB85EB CRC64;

Query Match 81.8%; Score 36; D5 1; Length 300;
Best Local Similarity 75.0%; Pred. No. 16; Mismatches 0; Indels 0; Gaps 0;
Matches 6; Conservative 2;

QY 1 ILGCIITS 8
DB 133 ILGCELS 140

RESULT 4
EGLN_PIG STANDARD; PRT: 653 AA.
AC P37176;
DT 01-OCT-1994 (Rel. 30, Created)
DT 01-OCT-1994 (Rel. 30, Last sequence update)
DI 28-FEB-2003 (Rel. 41, Last annotation update)
DE Endoglin precursor.
EN ENG.
OS Sus scrofa (Pig).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Suidae; Suidae.
OX NCBI_TaxID=9823;
RN (1)
RP SEQUENCE FROM N.A.
RC TISSUE-Uterus;
RX MEDLINE=94124550; PubMed=8294451;
RA Yamashita H., Ichijo H., Grimsby S., Moren A., ten Dijke P.,
RA Miyazono K.;
RT "Endoglin forms a heteromeric complex with the signaling receptors
RT for transforming growth factor-beta.";
RL J. Biol. Chem. 269:1995-2001(1994).
CC -!- FUNCTION: MAJOR GLYCOPROTEIN OF VASCULAR ENDOTHELIUM. MAY PLAY A
CC CRITICAL ROLE IN THE BINDING OF ENDOTHELIAL CELLS TO INTEGRINS
CC AND/OR OTHER RGD RECEPTORS.
CC -!- SUBUNIT: HOMODIMER THAT FORMS AN HETEROMERIC COMPLEX WITH THE
CC SIGNALING RECEPTORS FOR TRANSFORMING GROWTH FACTOR-BETA: TGF-BETA
CC RECEPTORS 1 AND/OR II. IT IS ABLE TO BIND TGF-BETA 1, AND 3
CC EFFICIENTLY AND TGF-BETA 2 LESS EFFICIENTLY.
CC -!- SUBCELLULAR LOCATION: Type I membrane protein.
CC -!- SIMILARITY: SOME REGIONS OF SIMILARITY TO TGF-BETA RECEPTOR TYPE
CC III (BETAGLYCAN).
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@sib-sib.ch).
CC
CC EMBL: Z23142; CAA00679.1;
CC PIR: A49722; A49722.
CC InterPro: IPR001507; Endoglin/CD105.
CC Pfam: PF00100; zora_pellucida; 1.
CC SMART: SM00241; 2P; 1.
CC Cell adhesion; Glycoprotein; Transmembrane; Signal.
CC SIGNAL 1 24 POTENTIAL.
CC CHAIN 25 653 ENDOLIN.
CC DOMAIN 25 581 EXTRACELLULAR (POTENTIAL).
CC TRANSMEM 582 606 POTENTIAL.

```

```

FT DOMAIN 607 653 CYTOPLASMIC (POTENTIAL).
FT DOMAIN 335 574 SER/THR-RICH.
FT SITE 396 400 CELL ATTACHMENT SITE (BY SIMILARITY).
FT CARBOHYD 57 57 N-LINKED (GLCNAC...) (POTENTIAL).
FT CARBOHYD 306 306 N-LINKED (GLCNAC...) (POTENTIAL).
SQ SEQUENCE 653 AA: 70279 MW: 7887B4A61DFE3E5A CRC64;

Query Match 79.5%; Score 35; D5 1; Length 653;
Best Local Similarity 87.5%; Pred. No. 46; Mismatches 0; Indels 0; Gaps 0;
Matches 7; Conservative 0;

QY 2 LGCIITS 9
DB 379 LGCIITS 386

RESULT 5
YBT6_YEAST STANDARD; PRT: 946 AA.
AC P36250;
DT 01-OCT-1994 (Rel. 30, Created)
DT 01-OCT-1994 (Rel. 30, Last sequence update)
DT 01-NOV-1995 (Rel. 32, Last annotation update)
DE Hypothetical 105.9 kDa protein in AAC3-REF5 intergenic region.
GN YBC86C OR YBR0809.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomycos.
OX NCBI_TaxID=4932;
RN (1)
RP SEQUENCE FROM N.A.
RC STRAN-S288C;
RX MEDLINE=95208357; PubMed=7900426;
RA Mannhaupt G., Stucka R., Ehmele S., Vetter I., Feldmann H.;
RT "Analysis of a 70 kb region on the right arm of yeast chromosome II.";
RL Yeast 10:1363-1381(1994).
CC -!- SUBCELLULAR LOCATION: Integral membrane protein (Potential).
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@sib-sib.ch).
CC
CC EMBL: X78993; CAA55593.1;
CC EMBL: Z35955; CAA85034.1;
CC PIR: S48255; S48255.
CC SGD: S0000290; IST2.
CC Pfam: PF04547; DUF590; 1.
CC Hypothetical protein; Transmembrane.
CC TRANSMEM 122 142 POTENTIAL.
CC TRANSMEM 154 174 POTENTIAL.
CC TRANSMEM 218 238 POTENTIAL.
CC TRANSMEM 254 274 POTENTIAL.
CC TRANSMEM 303 323 POTENTIAL.
CC TRANSMEM 448 468 POTENTIAL.
CC TRANSMEM 506 526 POTENTIAL.
CC TRANSMEM 564 584 POTENTIAL.
CC SEQUENCE 946 AA: 105903 MW: F51A43A5D378B7BC CRC64;

Query Match 77.3%; Score 34; D5 1; Length 946;
Best Local Similarity 62.5%; Pred. No. 96; Mismatches 3; Conservative 0; Indels 0; Gaps 0;
Matches 5;

QY 1 ILGCIITS 8
DB 520 ILGCVITA 527

RESULT 6
T2R7_HUMAN

```


RA Fahey J., Helton E., Kettiman M., Madan A., Rodrigues S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Zoufard G.G.,
 RA Blakesley R.W., Grisham J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Trifunovic J., Schmutz J., Myers R.M.,
 RA Butterfield V.S.N., Krzyzinski M.I., Skalska U., Smallus D.E.,
 RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.,
 RT *Generation and initial analysis of more than 15,000 full-length
 human and mouse cDNA sequences.*;
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [4]
 RN STRUCTURE BY NMR OF 1-105.
 RX MEDLINE-9426557; PubMed-8208296;
 RA Yoon H.S., Hajduk P.J., Petros A.M., Olejniczak E.T., Meadows R.P.,
 RA Fesik S.W.,
 RT *Solution structure of a pleckstrin-homology domain.*;
 RL Nature 369:672-675(1994).
 CC EXACT FUNCTION: MAJOR PROTEIN KINASE C SUBSTRATE OF PLATELETS. ITS
 CC -!- FUNCTION: Contains 1 DEP domain.
 CC -!- SIMILARITY: Contains 2 PH domains.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL; X07743; CAA00564.1; -;
 DR EMBL; H018549; AAI18549.1; -;
 DR PIR; S00755; S00755.
 DR PDB; 1PLS; 03-JUN-95.
 DR Genew; HGNC:9070; PLEK.
 DR MIM; 173570; -;
 DR GO; GO:0005509; Focal-calcium ion binding activity; TAS.
 DR InterPro; IPR000591; DEP.
 DR Pfam; PF00610; IPR001849; PH.
 DR Pfam; PF00169; PH; 2.
 DR SMART; SM00049; DEP; 1.
 DR SMART; SM00233; PH; 2.
 DR PROSITE; PS0186; DEP; 1.
 DR PROSITE; PS50003; PH; DOMAIN; 2.
 KW Phosphorylation; Repeat; 3D-structure.
 FT DOMAIN 4 101
 FT DEP 1.
 FT DOMAIN 136 221
 FT PH 2.
 FT VARIANT 92 92
 FT W-5; R.
 FT /FTID-VAR_005524.
 FT N -> K (IN REF. 3).
 FT CONFLICT 97 97
 FT STRAND 7 14
 FT STRAND 22 29
 FT TURN 30 31
 FT STRAND 32 36
 FT TURN 39 40
 FT TURN 40 40
 FT STRAND 46 49
 FT STRAND 55 56
 FT STRAND 58 73
 FT TURN 74 76
 FT STRAND 77 82
 FT HELIX 86 103
 FT TURN 104 105
 SQ SEQUENCE 350 AA; 40082 MW; 2E2A128CB526361 CRC64;
 Query Match 75.0%; Score 33; DB 1; Length 350;
 Best Local Similarity 55.6%; Pred. No. 63;
 Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
 QY 1 LLGC:ITSL 9
 DB 292 LRCCVTVSV 300

RESULT 8
 PLEK_MOUSE STANDARD; PRI; 350 AA.
 ID PLEK_MOUSE
 AC O9JHK5; O9FRI9;
 DT 28-FEB-2003 (Rel. 41, Created)
 DI 28-FEB-2003 (Rel. 41, Last sequence update)
 DI 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Pleckstrin.
 GN PLEK.
 OS Mus musculus (Mouse).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 CX NCBI_TaxID=10090;
 RN [1]
 RN SEQUENCE FROM N.A.
 RC STRAIN=BALE/C;
 RX MEDLINE-2031822; PubMed-10860665;
 RA Omarik J.L., Hegamyer G., Gerrard B., Dean M., Colburn N.H.,
 RT "cDNA cloning and mapping of mouse pleckstrin (Plek), a gene
 RI upregulated in transformation-resistant cells.*;
 RL Genomics 66:204-212(2000).
 RN [2]
 RN SEQUENCE FROM N.A.
 RA Zhang Y., Wu G., Paige C.J.,
 RT "Involvement of pleckstrin in B cell differentiation and
 RT activation.*;
 RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RN SEQUENCE FROM N.A.
 RA Ann H.-J., Cho J.-J.,
 FT "Mouse pleckstrin-1 is induced in mast cells after IgE cross-
 RT linking.*;
 RL Submitted (SEP-2000) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: MAJOR PROTEIN KINASE C SUBSTRATE OF PLATELETS, ITS
 CC EXACT FUNCTION IS NOT KNOWN.
 CC -!- SIMILARITY: Contains 1 DEP domain.
 CC -!- SIMILARITY: Contains 2 PH domains.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL; AF181829; AAF75830.1; -;
 DR EMBL; AF673294; AAF72039.1; -;
 DR EMBL; AF303745; AAG29513.1; -;
 DR HSSP; P04567; IPI5.
 DR MGD; MGI:1860485; Plek.
 DR InterPro; IPR000591; DEP.
 DR InterPro; IPR001849; PH.
 DR Pfam; PF00610; DEP; 1.
 DR Pfam; PF00169; PH; 2.
 DR SMART; SM00049; DEP; 1.
 DR SMART; SM00233; PH; 2.
 DR PROSITE; PS0186; DEP; 1.
 DR PROSITE; PS50003; PH; DOMAIN; 2.
 KW Phosphorylation; Repeat.
 FT DOMAIN 4 101
 FT DEP 1.
 FT DOMAIN 136 221
 FT PH 2.
 FT CONFLICT 120 120 L -> Q (IN REF. 3).
 FT CONFLICT 225 225 F -> L (IN REF. 3).
 SQ SEQUENCE 350 AA; 39900 MW; 348F3CB469B7CC53 CRC64;
 Query Match 75.0%; Score 33; DB 1; Length 350;
 Best Local Similarity 55.6%; Pred. No. 63;
 Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
 QY 1 LLGC:ITSL 9

```

Db          292 LAGCWTISV 300
! ! ! ! !
RESULT 9
ARGD_PSEPK STANFORD: PRT: 406 AA.
AC P59319;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Acetylornithine aminotransferase (EC 2.6.1.11) (NCOR1).
GN ARGD OR PP4481.
OS Pseudomonas putida (strain KT2440).
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
OC Pseudomonadaceae; Pseudomonas.
OX NCBI_TaxID=160488;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22423060; PubMed=12534463;
RA Nelson K.E., Weinel C., Paulsen I.T., Dodson R.J., Hilbert H.,
RA Martins dos Santos V.A.P., Feuts D.E., Gill S.R., Pop M., Holmes M.,
RA Brinkac C., Beanan M., Deacy R.T., Daugherty S., Conway J.,
RA Madupu K., Nelson W., White O., Peterson J., Khouri H., Hanco I.,
RA Chris Lee P., Holtzapple E., Scanlan D., Tran K., Motzke A.,
RA Utterback T., Rizzo M., Lee K., Kosack D., Moestl D., Wedler H.,
RA Lauber J., Stjepandic D., Hebeisel J., Straetz M., Heim S.,
RA Kiewitz C., Eisen J., Tirmis K.N., Duesterhoeft A., Tiedler B.,
RA Fraser C.W.;
RI "Complete genome sequence and comparative analysis of the
RI metabolically versatile Pseudomonas putida KT2440."
RL Environ. Microbiol. 4:799-808(2002).
CC -1- CATALYTIC ACTIVITY: N(2)-acetyl-L-ornithine + 2-oxoglutarate ->
CC acetyl-L-glutamate + 5-semialdehyde + L-glutamate.
CC -1- COFACTOR: Pyridoxal phosphate (by similarity).
CC -1- PATHWAY: Arginine biosynthesis; fourth step.
CC -1- MISCELLANEOUS: May also have succinyl-L-aminopimelate
CC aminotransferase activity, thus carrying out the fourth step in
CC lysine biosynthesis.
CC -1- SIMILARITY: Belongs to class-II; of pyridoxal phosphate-dependent
CC aminotransferases.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement. (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC
CC EMBL: AEC16760; AAN70056.1; -
CC IIGR: PP4481; -
CC HAMAP: MF_01107; -1.
CC Pfam: PF00202; aminotran_3; 1.
CC TIGRfams: TIGR00707; argD; 1.
CC PROSITE: PS00600; AA_TRANSFER_CLASS_3; 1.
CC Arginine biosynthesis; Transferase; Aminotransferase;
KW Pyridoxal phosphate; Complete proteome.
FT BINDING 255 255 PYRIDOXAL PHOSPHATE (BY SIMILARITY).
SQ SEQUENCE 406 AA; 43490 MW; 5597D2F10C3E221E CRC64;
Query Match 75.0%; Score 33; DB 1; Length 406;
Best Local Similarity 71.4%; Pred. No. 72;
Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
QY 1 LLGCVLT 7
Db 341 LLGCVLT 347
! ! ! ! !
RESULT 10
FLHF_PSEPU STANFORD: PRT: 437 AA.
ID FLHF_PSEPU

```

```

AC 052256;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE Flagellar biosynthesis protein flhF (Flagella associated GTP-binding
DE protein).
GN FLHF.
OS Pseudomonas putida.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
OC Pseudomonadaceae; Pseudomonas.
OX NCBI_TaxID=303;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=PRS2000;
RX MEDLINE=98164369; PubMed=9503621;
RA Ditty J.L., Grimm A.C., Harwood C.S.;
RT Identification of a chemotaxis gene region from Pseudomonas putida.;
RL FEMS Microbiol. Lett. 159:267-273(1998).
CC -1- FUNCTION: NECESSARY FOR FLAGELLUM BIOSYNTHESIS. MAY BE INVOLVED
CC IN TRANSLOCATION OF THE FLAGELLUM (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE SRP FAMILY OF GTP-BINDING PROTEINS.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement. (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC
CC EMBL: AF031898; AAC08059.1; -
CC InterPro: IPR003593; AAA_Arase.
CC InterPro: IPR008957; SRP54.
CC Pfam: PF00448; SRP54; 1.
CC Pfam: PF02381; SRP54; 1.
CC Pfam: PF02381; SRP54; 1.
CC Pfam: PF02381; SRP54; 1.
CC PRODOM: P0000819; SRP54; 1.
CC SMART: SM00382; AAA; 1.
CC PROSITE: PS00300; SRP54; FALSE NEG.
CC Flagella: GTP-binding.
KW NP_BIND 225 232 GTP (BY SIMILARITY).
FT NP_BIND 303 307 GTP (BY SIMILARITY).
FT NP_BIND 362 364 GTP (BY SIMILARITY).
SQ SEQUENCE 437 AA; 47512 MW; EBFA6A650B9B27A CRC64;
Query Match 75.0%; Score 33; DB 1; Length 437;
Best Local Similarity 66.7%; Pred. No. 76;
Matches 6; Conservative 1; Mismatches 2; Indels 0; Gaps 0;
QY 1 LLGCVLTSL 9
Db 355 LLGCVLTSL 363
! ! ! ! !
RESULT 11
YBSE_BACSU STANFORD: PRT: 441 AA.
ID YBSE_BACSU
AC 005213; O08069; Q45578;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Hypothetical protein ybde precursor.
GN ybde.
OS Bacillus subtilis.
OC Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus.
OX NCBI_TaxID=1423;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=168;
RX MEDLINE=97419514; PubMed=9274029;
RA Liu H., Hage K., Yasumoto K., Ohashi Y., Yoshikawa H., Takahashi H.;
RT "Sequence and analysis of a 31 kb segment of the Bacillus subtilis
RT chromosome in the area of the rnh and rmg operons."
RL Microbiology 143:2763-2767(1997).

```



```

Db          591 MLCQVIT 597
|||||
RESULT 13
PGDS_HUMAN  STANDARD:      PRT: 1089 AA.
AC  P16234;
DI  01-APR-1390 (Rel. 14, Last sequence update)
DT  01-APR-1390 (Rel. 14, Last sequence update)
DT  15-SEP-2003 (Rel. 42, Last annotation update)
DE  Alpha platelet-derived growth factor receptor precursor (EC 2.7.1.112)
DE  (PDGFR-R-alpha) (CD140a antigen).
CN  PDGFRA.
OS  Homo sapiens (Human).
OC  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC  Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
CX  NCBI_TaxID=9606;
[1]
RN  SEQUENCE FROM N.A.
RP  MEDLINE=89130149; PubMed=2536956;
RA  Matsui T., Heidaran M., Miki T., Popescu N., la Rochelle W.,
RA  Kraus M., Pierce J., Aaronson S.;
RT  Isolation of a novel receptor cDNA establishes the existence of two
RT  PDGF receptor genes.;
RL  Science 243:800-804(1989).
[2]
RN  SEQUENCE FROM N.A.
RP  MEDLINE=89296915; PubMed=2544881;
RA  Claesson-Welsh L., Eriksson A., Westermark B., Heidin G.H.;
RT  cDNA cloning and expression of the human A-type platelet-derived
RT  growth factor (PDGF) receptor establishes structural similarity to
RT  the B-type PDGF receptor.;
RL  Proc. Natl. Acad. Sci. U.S.A. 86:4917-4921(1989).
[3]
RN  SEQUENCE FROM N.A.
RP  TISSUE-H-006;
RA  Kawagishi J., Ku T.;
RT  "Structure, Organization, and transcription units of the human
RT  alpha-platelet-derived growth factor receptor gene, PDGFRA.";
RL  Genomics 30:224-233(1995);
CC  -!- FUNCTION: THIS RECEPTOR BINDS PLATELET-DERIVED GROWTH FACTOR AND
CC  HAS A TYROSINE-PROTEIN KINASE ACTIVITY. THIS RECEPTOR CAN BIND
CC  EITHER PDGF-A OR PDGF-B.
CC  -!- CATALYTIC ACTIVITY: ATP + a protein:tyrosine -> ADP + protein
CC  tyrosine phosphate.
CC  -!- SUBUNIT: DIMER OF EITHER ALPHA-ALPHA, BETA-BETA OR ALPHA-BETA
CC  SUBUNITS.
CC  -!- SUBCELLULAR LOCATION: Type I membrane protein.
CC  -!- SIMILARITY: BELONGS TO THE CSF-1/PDGF RECEPTOR FAMILY OF TYROSINE-
CC  PROTEIN KINASES.
CC  -!- SIMILARITY: Contains 5 immunoglobulin-like c2-type domains.
CC  -----
CC  THIS SWISS-PROT entry is copyright. It is produced through a collaboration
CC  between the Swiss Institute of Bioinformatics and the EMBL Outstation:
CC  the European Bioinformatics Institute. There are no restrictions on its
CC  use by non-profit institutions as long as its content is in no way
CC  modified and this statement is not removed. Usage by and for commercial
CC  entities requires a license agreement (See http://www.isb-sib.ch/announcement/
CC  or send an email to license@isb-sib.ch).
CC  -----
DR  EMBL; M21574; AAA96715.1; -
DR  EMBL; M22734; AAA60048.1; -
DR  EMBL; D50017; BAA08742.1; -
DR  EMBL; D50302; BAA08742.1; JOINED.
DR  EMBL; D50003; BAA08742.1; JOINED.
DR  EMBL; D50004; BAA08742.1; JOINED.
DR  EMBL; D50005; BAA08742.1; JOINED.
DR  EMBL; D50006; BAA08742.1; JOINED.
DR  EMBL; D50007; BAA08742.1; JOINED.
DR  EMBL; D50008; BAA08742.1; JOINED.
DR  EMBL; D50009; BAA08742.1; JOINED.

```

```

DR  EMBL; D50010; BAA08742.1; JOINED.
DR  EMBL; D50011; BAA08742.1; JOINED.
DR  EMBL; D50012; BAA08742.1; JOINED.
DR  EMBL; D50013; BAA08742.1; JOINED.
DR  EMBL; D50014; BAA08742.1; JOINED.
DR  EMBL; D50015; BAA08742.1; JOINED.
DR  EMBL; D50016; BAA08742.1; JOINED.
DR  PIR; A40162; PFHUGA.
DR  HSSP; P11362; IFCK.
DR  Genew; HGNC:8803; PDGFRA.
DR  MIM; 173490; -
DR  GO; GO:0005887; C: integral to plasma membrane; TAS.
DR  GO; GO:0005018; F: platelet-derived growth factor, alpha-recep. .; TAS.
DR  GO; GO:0008283; P: cell proliferation; TAS.
DR  GO; GO:0007166; P: cell surface receptor linked signal transdu. .; TAS.
DR  InterPro; IPR007110; Ig-Like.
DR  InterPro; IPR003598; Ig_C2.
DR  InterPro; IPR003006; Ig_MHC.
DR  InterPro; IPR000719; Prot_Kinase.
DR  InterPro; IPR001324; RTKinase11.
DR  InterPro; IPR001245; Tyr_pkinase.
DR  Pfam; PF00347; Ig_3.
DR  Pfam; PF00069; pkinase; 1.
DR  ProDom; PD000001; Prot_Kinase; 2.
DR  SMART; SM00408; IgC2; 1.
DR  SMART; SM00219; TyKc; 1.
DR  PROSITE; PS00835; IG_LIKE; 2.
DR  PROSITE; PS00107; PROTEIN_KINASE_ATP; 1.
DR  PROSITE; PS00111; PROTEIN_KINASE_DOM; 1.
DR  PROSITE; PS00109; PROTEIN_KINASE_TYR; 1.
DR  PROSITE; PS00240; RECEPTOR_TYR_KIN_III; 1.
KW  Tyrosine-protein kinase; Receptor; Transmembrane; Glycoprotein;
KW  Transferrase; Phosphorylation; ATP-binding; Immunoglobulin domain;
KW  Signal; Repeat.
FT  SIGNAL 1 23
FT  CHAIN 24 1089
FT  DOMAIN 24 524
FT  TRANSMEM 525 549
FT  DOMAIN 550 1089
FT  DOMAIN 24 113
FT  DOMAIN 202 306
FT  DOMAIN 319 430
FT  DOMAIN 593 954
FT  DOMAIN 1041 1087
FT  NE_BIND 599 607
FT  BINDING 627 627
FT  ACT_SITE 818 818
FT  MOD_RES 849 849
FT  CARBOHYD 42 42
FT  CARBOHYD 76 76
FT  CARBOHYD 103 103
FT  CARBOHYD 179 179
FT  CARBOHYD 353 353
FT  CARBOHYD 359 359
FT  CARBOHYD 458 458
FT  CARBOHYD 468 468
SQ  SEQUENCE 1089 AA: 122669 MW: 5E3FB9940ACD1BE8 CRC64;
Query Match 75.0%; Score 33; DB 1; Length 1089;
Best Local Similarity 55.6%; Pred. No. 1.6e-02;
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
QY 1 LIGCIITSL 9
DB 10 VLGCLLTGL 18
RESULT 14
DICE_MOUSE
ID DICE_MOUSE STANDARD: PRT: 1906 AA.
AC Q8R418;
DT 28-FEB-2003 (Rel. 41, Created)

```

DI 28-FEB-2003 (Rel. 41, last sequence update)
 DT 28-FEB-2003 (Rel. 41, last sequence update)
 DE Endoribonuclease Dicer (EC 3.1.26.-) (Double-strand-specific
 DE ribonuclease -Dicer-1)
 GN DICER1 OR DICER OR MDCR.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10030;
 RN [1]
 RP MEDLINE=21886641; PubMed=11889553;
 RX Nicholson R.H., Nicholson A.W.
 RA "Molecular characterization of a mouse cDNA encoding Dicer, a
 RI ribonuclease III ortholog involved in RNA interference."
 RL Mamm. Genome 13:67-73(2002).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX STRAIN=Czech II;
 RA Stroboda P., Andor M., Stein P., Schultz R.M.,
 RL "Mouse dicer homolog in oocyte and preimplantation embryos."
 RN Submitted (OCT-2001) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RA Doi N., Zenno S., Ui-Tei K., Takahashi F., Ueda R., Miyata Y.,
 RA Saigo K.
 RA "EIF2C family proteins and Dicer homologs are required for siRNA
 RT mediated RNAi in mammalian cells."
 RL Submitted (MAR-2002) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: Involved in cleaving double-stranded RNA in the RNA
 CC interference (RNAi) pathway. It produces 21 to 23 bp dsRNAs
 CC (siRNAs) which target the selective destruction of homologous
 CC RNAs.
 CC -!- TISSUE SPECIFICITY: Expressed in a wide variety of tissues.
 CC -!- SIMILARITY: BELONGS TO THE HELICASE FAMILY.
 CC -!- SIMILARITY: Contains 2 RNase III domains.
 CC -!- SIMILARITY: Contains 1 PAZ domain.
 CC -!- SIMILARITY: Contains 1 DRBM (double-stranded RNA-binding) domain.
 CC This SWISS-PROT entry is copyrighted. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@sib-sib.ch).
 CC -----
 DR EMBL: AF484524; AAL84636.1;
 DR EMBL: AF430845; AAM21495.1; AIT_LINT.
 DR EMBL: AB061470; BAC15765.1;
 DR MGD: MGI:217179; Dicer1.
 DR InterPro: IPR001410; DEAD.
 DR InterPro: IPR011159; DS_RBD.
 DR InterPro: IPR035034; NUF283.
 DR InterPro: IPR001650; Helicase_C.
 DR InterPro: IPR003100; PAZ.
 DR InterPro: IPR000099; RNase_3.
 DR Pfam: PF00270; DEAD_1.
 DR Pfam: PF00035; dsrm; 1.
 DR Pfam: PF03368; DUF283; 1.
 DR Pfam: PF00271; helicase_C; 1.
 DR Pfam: PF02170; PAZ; 1.
 DR Pfam: PF00636; Ribonuclease_3; 2.
 DR SMART: SM00487; DEXDC; 1.
 DR SMART: SM00359; DSRM; 1.
 DR SMART: SM00490; HELIC_C; 1.
 DR SMART: SM00535; RIBOC; 2.
 DR PROSITE: PS50137; DS_RBD; 1.
 DR PROSITE: PS50821; PAZ; 1.
 DR PROSITE: PS00517; RNASE_3_1; 1.
 DR PROSITE: PS50142; RNASE_3_2; 2.
 KW Helicase; ATP-binding; RNA-binding; HydroLase; Nuclease; Endonuclease;
 KW Repeat.

FT NP_HIND 34 42 ATP (POTENTIAL).
 FT SITE 165 168 DECH BOX.
 FT DOMAIN 881 1032 PAZ.
 FT DOMAIN 1266 1393 RNASE III 1.
 FT DOMAIN 1650 1808 RNASE III 2.
 FT DOMAIN 1833 1898 DRBM.
 FT CONFLICT 97 97 A -> C (IN REF. 2).
 FT CONFLICT 157 157 S -> P (IN REF. 3).
 FT CONFLICT 279 279 H -> Y (IN REF. 2).
 FT CONFLICT 600 600 A -> T (IN REF. 3).
 FT CONFLICT 749 749 E -> D (IN REF. 2).
 FT CONFLICT 827 827 T -> I (IN REF. 2).
 FT CONFLICT 878 878 G -> S (IN REF. 1).
 FT CONFLICT 955 955 Y -> C (IN REF. 3).
 FT CONFLICT 983 983 I -> A (IN REF. 3).
 FT CONFLICT 1080 1080 R -> G (IN REF. 3).
 FT CONFLICT 1100 1100 T -> S (IN REF. 3).
 FT CONFLICT 1326 1326 P -> H (IN REF. 1).
 FT CONFLICT 1609 1609 A -> S (IN REF. 3).
 FT CONFLICT 1850 1850 K -> E (IN REF. 3).
 SQ SEQUENCE 1906 AA: 215756 MW: 230EA9BFC19F3091 CRC64:
 Query Match 75.0% Score 33; DB 1; Length 1906;
 Best local Similarity 75.0% Pred. No. 2.6e+02;
 Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 QY 1 LLGCITTS 8
 DB 1552 LLGCYLTTS 1559
 RESULT 15
 DICE_HUMAN
 ID DICE_HUMAN STANDARD; PRT: 1912 AA.
 AC Q9UPF3; O95943; Q9UQ02;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Endoribonuclease Dicer (EC 3.1.26.-) (Helicase with RNase motif)
 DE (Helicase-MOI).
 GN DICER1 OR DICER OR HERNA OR KIAA0928.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=20246304; PubMed=10786632;
 RA Matsuda S., Ichigotani Y., Okuda T., Irimura I., Nakatsugawa S.,
 RA Hamaguchi M.;
 RT "Molecular cloning and characterization of a novel human gene (HFRNA)
 RT which encodes a putative RNA-helicase."
 RL Biochim. Biophys. Acta 1490:163-169(2000).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Lung;
 RA Provost P., Dishart D., Doucet D., Hermanson A., Frendewey D.,
 RA Samuelsson B., Radmark O.;
 RT "RNA binding and processing by recombinant human Dicer."
 RL Submitted (JUN-2002) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Brain;
 RX MEDLINE=99246063; PubMed=10211032;
 RA Nagase T., Ishikawa K.-I., Suyama M., Kikuno R., Hirose M.,
 RA Miyajima N., Tanaka A., Kotani H., Nomura N., Ohara O.;
 RT "Prediction of the coding sequences of unidentified human genes. XII.
 RT The complete sequences of 100 new cDNA clones from brain which code
 RT for large proteins in vitro."
 RL DNA Res. 6:63-70(1999).
 RN [4]
 RP REVISIONS.
 RX MEDLINE=22158633; PubMed=12168954;

RA Nakajima D., Okazaki N., Yarakawa H., Kikuno A., Ohara O., Nagase T.;
RT "Construction of expression-ready cDNA clones for KIAA genes: manual
RL curation of 330 KIAA cDNA clones.";
RL DNA Res. 9:99-106(2002).
RN [5]
RP SEQUENCE OF 1238-1912 FROM N.A.
RC TISSUE=Lung;
RX MEDLINE=99162526; PubMed=10051563;
RA Provost P., Samuelsson A., Kadmark O.;
RI "Interaction of 5-lipoxygenase with cellular proteins.";
RL Proc. Natl. Acad. Sci. U.S.A. 96:1881-1885(1999).
CC -!- FUNCTION: Involved in cleaving double-stranded RNA in the RNA
CC interference (RNAi) pathway. It produces 21 to 23 bp dsRNAs
CC (siRNAs) which target the selective destruction of homologous
CC RNAs.
CC -!- SIMILARITY: BELONGS TO THE HELICASE FAMILY.
CC -!- SIMILARITY: Contains 2 RNase III domains.
CC -!- SIMILARITY: Contains 1 PAZ domain.
CC -!- SIMILARITY: Contains 1 DRRM (double-stranded RNA-binding) domain.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (see <http://www.isg-sib.ch/announc/>
CC or send an email to license@sib-sib.ch).
CC
DR EMBL: AR028449; BAA78691.1; ALT_INIT.
DR EMBL: A313226.1; CAB38857.2; ..
DR EMBL: ABC23145; BAA76772.2; ALT_INIT.
DR Genew: HGNC:17098; DICER1.
DR MIM: 606241; ..
DR InterPro: IPR001410; DEAD.
DR InterPro: IPR001159; DS_RBD.
DR InterPro: IPR005034; DUF283.
DR InterPro: IPR001650; Helicase_C.
DR InterPro: IPR003100; PAZ.
DR InterPro: IPR006999; RNase_3.
DR Pfam: PF00270; DEAD_1.
DR Pfam: PF00635; Gsmr_1.
DR Pfam: PF03368; DUF283; 1.
DR Pfam: PF00271; Helicase_C; 1.
DR Pfam: PF02170; PAZ; 1.
DR Pfam: PF00636; Ribonuclease_3; 2.
DR SMART: SM00487; DEXDC; 1.
DR SMART: SM00358; DSRM; 1.
DR SMART: SM00490; HELIC_C; 1.
DR SMART: SM00535; RIBOC; 2.
DR PROSITE: PS0137; DS_RBD; 1.
DR PROSITE: PS0821; PAZ; 1.
DR PROSITE: PS0517; RNase_3; 1; 1.
DR PROSITE: PS0142; RNase_3; 2; 2.
KW Helicase; ATP-binding; RNA-binding; Hydrolyase; Nuclease; Endonuclease;
KW Repeat.
FT NP_BIND 34 41 ATP (POTENTIAL).
FT SITE 165 168 DECH_BOX.
FT DOMAIN 88 1632 PAZ.
FT DOMAIN 1266 1393 RNase III 1.
FT DOMAIN 1656 1814 RNase III 2.
FT DOMAIN 1839 1904 DRRM.
FT CONFLICT 65 80 VIKTKELSYQIRGDFS -> STLLKCYLDAGETSA
(IN REF. 1).
FT CONFLICT 179 179 I -> F (IN REF. 1).
FT CONFLICT 185 185 N -> Z (IN REF. 1).
FT CONFLICT 204 204 C -> W (IN REF. 1).
FT CONFLICT 208 208 E -> D (IN REF. 1).
FT CONFLICT 213 213 I -> F (IN REF. 1).
FT CONFLICT 363 364 QQ -> HS (IN REF. 1).
FT CONFLICT 382 384 KO -> NT (IN REF. 1).
FT CONFLICT 482 483 D -> H (IN REF. 1).
FT CONFLICT 599 599 ..
SQ SEQUENCE 1912 AA: 217627 MW: 996399984807421 CRC64;

Query Match 75.0%; Score 33; DR 1; Length 1912;
Best Local Similarity 75.0%; Pred. No. 2.6e+02;
Matches 6; Conservative 1; Mismatches 0; Gaps 0;
QY 1 LLGCITTS 8
Db 1556 LLGCYLTTS 1563
|||||
|
|
|
|
|

Search completed: September 29, 2003, 19:07:47
Job time : 13 secs

GenCore version 5.1.4
Copyright (c) 1993 - 2003 CompuGen Ltd.

QM protein - nucleic search, using frame_plus_p2c model

Run on: September 30, 2003, 10:34:19 : Search time 2202 Seconds
(without alignments)
167.206 Million cell updates/sec

Title: US-09-965-594-1_COPY_14_22
Perfect score: 44
Sequence: 1 LLAGITSL 9

Scoring table: BL0SUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Dgapop 6.0 , Delext 7.0

Searched: 268871 seqs, 2045481386 residues

Total number of hits satisfying chosen parameters: 5776890

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 99%
Listing first 45 summaries

Command line parameters:

-MODEL=frame_plus_p2c_model -DEV_X11
-Q/cqn211/USPIC_spool/US09965594/runat_29092003_164249_14072/app_query.fasta_1.199
-DB=GenEmbl -Qfmt=fastap -SUFFIX=gc -MINMA=CH-0.1 -DOOP=0 -DOOPEXT=0
-UNITS=bits -START=1 -END=1 -MATRIX=blosum62 -TRANS=human40.cdi -DIST=45
-DOCALIGN=200 -THR_SCORE=pct -THR_MAX=99.9 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=ptc -NOR=ext -HEAPSIZE=1000 -MINLEN=0 -MAXLEN=200000000
-USER=US09965594 -CGN_1_1_3508 -runat_29092003_164249_14072 -NCPU=6 -ICPU=3
-NO_MMAP -LARGEQUERY -NGI_SCORES=0 -WAIT -DSPBCHECK=100 -LONGLOG
-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -EFLOP=6 -DELEXT=7

Database : GenBank

1: gb_kat:
2: gb_hsq:
3: gb_tet:
4: gb_ov:
5: gb_ov:
6: gb_pat:
7: gb_ph:
8: gb_pl:
9: gb_pri:
10: gb_ro:
11: gb_sts:
12: gb_sy:
13: gb_un:
14: gb_vi:
15: em_ba:
16: em_fus:
17: em_hum:
18: em_ie:
19: em_mu:
20: em_or:
21: em_or:
22: em_ov:
23: em_pat:
24: em_ph:
25: em_pl:
26: em_ro:
27: em_sts:
28: em_un:

29: em_vit:
30: em_hsq_hum:
31: em_hsq_inv:
32: em_hsq_other:
33: em_hsq_mus:
34: em_hsq_pln:
35: em_hsq_ror:
36: em_hsq_mam:
37: em_hsq_vit:
38: em_sy:
39: em_hsq_hum:
40: em_hsq_mus:
41: em_hsq_other:

pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	43	97.7	543	14	AB072047 Hepatitis
2	43	97.7	543	14	AB072048 Hepatitis
3	43	97.7	543	14	AB072067 Hepatitis
4	43	97.7	540	14	AB072102 Hepatitis
5	43	97.7	540	14	AB089533 Hepatitis
6	43	97.7	540	14	AB089539 Hepatitis
7	43	97.7	540	14	AB089558 Hepatitis
8	43	97.7	540	14	AB100809 Hepatitis
9	43	97.7	540	14	AB100827 Hepatitis
10	43	97.7	1211	14	AY070171 Hepatitis
11	43	97.7	2691	14	AF187824 Cache Val
12	43	97.7	2691	14	AF231116 Cache Val
13	43	97.7	2691	14	AF231118 Cache Val
14	43	97.7	4463	14	AF186241 Cache Val
15	43	97.7	9359	14	AF13916 Hepatitis
16	43	97.7	9359	14	AF165053 Hepatitis
17	43	97.7	9379	14	AF165049 Hepatitis
18	43	97.7	9379	14	AF165050 Hepatitis
19	43	97.7	9434	14	HPCJTB
20	43	97.7	9436	6	E07266 Blood-sprea
21	43	97.7	9436	14	HPCJTA
22	43	97.7	9442	14	AY051292 Hepatitis
23	43	97.7	9598	14	AB049101 Hepatitis
24	43	97.7	217285	2	AC126317 Rattus no
25	43	97.7	260760	2	AC127861 Rattus no
26	42	95.5	543	14	AF169258 Hepatitis
27	42	95.5	819	6	E06893 DNA encodin
28	42	95.5	3461	6	E07544
29	42	95.5	3461	6	E09288
30	42	95.5	9375	14	AF207754 Hepatitis
31	42	95.5	9379	14	AF207768 Hepatitis
32	42	95.5	89352	9	AC006030 Homo sapi
33	42	95.5	105991	2	AC016960 Homo sapi
34	42	95.5	133801	9	AC034585 Homo sapi
35	42	95.5	138517	9	AC019288 Homo sapi
36	42	95.5	230396	2	AC103315 Rattus no
37	42	95.5	231275	2	AC099421 Rattus no
38	41	93.2	540	14	AB072056 Hepatitis
39	41	93.2	1096	14	AB013622 Hepatitis
40	41	93.2	8780	14	AF054253 Hepatitis
41	41	93.2	8780	14	AF054255 Hepatitis
42	41	93.2	8780	14	AF054259 Hepatitis
43	41	93.2	8781	14	AF054256 Hepatitis
44	41	93.2	9379	14	AF207773 Hepatitis
45	41	93.2	9448	14	HPCJ483

ALIGNMENTS

RESULT 1


```

DB
|||||.....
37 CTACTGGTGCATCGCTACTAGCCTC 63

RESULT 6
AB100809
LOCUS      540 bp      RNA      linear      VRL 02-APR-2003
DEFINITION Hepatitis C virus NS3 gene for polyprotein, partial cds, isolate:
            Y-CH-06-B.
ACCESSION AB100809
VERSION   AB100809.1 GI:29467656
KEYWORDS
SOURCE
ORGANISM   Hepatitis C virus
            Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
            Hepacivirus.

REFERENCE
AUTHORS    Ogata,S. and Hotta,H.
TITLE      Identification of Hepatitis C Virus (HCV) Subtype 1b Strains That
            Are Highly, or Only Weakly, Associated with Hepatocellular
            Carcinoma on the Basis of Secondary Structure of an Amino-Terminal
            Portion of the HCV NS3 Protein
JOURNAL    Unpublished
PUBLISHED
AUTHORS    Hotta,H.
TITLE      Direct Submission
JOURNAL    Submitted (24-JAN-2003) Hak Hotta, Kobe University Graduate School
            of Medicine, Division of Microbiology; Chuo-ku, Kusunoki-cho 7-5-1,
            Kobe, Hyogo 650-0017, Japan (E-mail:hotta@kobe-u.ac.jp,
            Tel:81-78-382-5500, Fax:81-78-382-5513)
            Location/Qualifiers
FEATURES   source
            1..540
                /organism="Hepatitis C virus"
                /mol_type="genomic RNA"
                /isolate="Y-CH-06-B"
                /db_xref="taxon:11103"
                /note="subtype: 1b"
            gene
            1..540
                /gene="NS3"
            CDS
            1..540
                /codon_start=1
                /product="polyprotein"
                /protein_id="BAC67219.1"
                /db_xref="GI:29467657"
                /translation="APITAYSHQIRGLGCVTSLTGRXKQVEGEVQVWVSTATGCSFL
                ATCVNGVQVTVHGASKTLAGPKGPIITQYTNVDGLVQWAPGARSMPTCTGSS
                DLVLYTRHADVIPVRHGGDGRGSLSPRPVSVLKSGSGPLFCPLGHVVVGIFRAAVCT
                RGVAKAVDFEVEVSEMETMR"
            BASE COUNT      93 a 171 c 166 g 110 t
            ORIGIN
            Alignment Scores:
            Pred. No.:      4.14      Length:      540
            Score:          43.00      Matches:      8
            Percent Similarity: 100.00%      Conservative: 1
            Best Local Similarity: 88.89%      Mismatches: 0
            Query Match:      97.73%      Indels:      0
            Db:              14      Gaps:      0
            US-09-965-594-1_COPY_14_22 (1-5) x AB100809 (1-540)

QY      1 LeuGugGlyCysIleIeThrSerLeu 9
|||||.....
DB      37 CTACTGGTGCATCGCTACTAGCCTC 63

RESULT 10
AB100809
LOCUS      1211 bp      RNA      linear      VRL 01-APR-2002
DEFINITION Hepatitis C virus strain 697 polyprotein gene, partial cds.
ACCESSION AB100809
VERSION   AB100809.1 GI:19879463
KEYWORDS
SOURCE
ORGANISM   Hepatitis C virus
            Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
            Hepacivirus.
            1 (bases 1 to 1211)
REFERENCE
AUTHORS    Kaimina,O., Norder,H., Mukomolov,S. and Magnus,L.O.
TITLE      A natural intergenotypic recombinant of hepatitis C virus
            identified in St. Petersburg
JOURNAL    J. Virol. 76 (8), 4034-4043 (2002)
MEDLINE    21904745
PUBMED     11907242
REFERENCE   2 (bases 1 to 1211)

```


AUTHORS Kalina, C., Nordcr, H., Mukomolov, S. and Magnus, L.
 TITLE Direct Submission
 JOURNAL Submitted (12-DEC-2001) Virology, Swedish Institute for Infectious
 Disease Control, Solna SE 171 82, Sweden

FEATURES
 Location/Qualifiers

1..1211
 /organism="Hepatitis C virus"
 /mol_type="genomic RNA"
 /strain="697"
 /db_xref="taxon:11103"
 /note="recombinant 2k/1b"
 1..1211
 /note="contains E2, p7, NS2 and NS3"
 /cddon_start=3
 /product="polyprotein"
 /protein_id="AAL58587.1"
 /db_xref="GI:19879464"

GDS
 /translation="ETDCTKKSLEIDC:GPMIDIGSCNKNTRSYDAEAQKLVQSQK
 ISQLDAPVSLGSSDAIKAIKRAQTYPTMHELETIFLMKNCQYTSFNHNSGISQ
 AKWLIKTIHFQICSRHSHTHPCRCINDGTQCGNMGWQFAGEMETYSKNDYFSD
 CMLFYTFNAPGITEFSFYLOLOKNTTGASKVLGLIKYGNMNFVGVWKTQGY
 LLLTYINTELSONRRLILATQFALSRSHSRQPSMSNAIOGSTTECHNAKGVGC
 ZSPWGPIGNLTACGDSFNKYIKTPEKLYSKNNKGVWCSNDVHCLNDFEPADREV
 VNKTKLCTLDVSVEDIFSVAS:CKMADGVCIVNCAKWNIIKDSGLIYFTDHR
 REGODTNDGFEYCISSCNTERFP:NPDIKADCTWEPHSRKSXYISILSDSBEFR
 RAMAKISHTLIYFNKPIANPLRPITYKFTANGVNSDGIKSAYLSSIPALGGI
 SDGVQSQSLFPLDMIVFDIAVIRTYVYHYDTGHTIGIYQHDHCTGPPAMV
 PHENWMTSOFRTSMGCFEFCCLAINTCGVFGSCOD:THPETKYRKAVEEVVLT
 VCLNFGNNACTE:NAIEBK:TDIEFLOKTVTITPLNLAQNHKLYSGQINDLGS
 FSGCGNVOKNHSILGKMTVAFQDYVCHGASRQVIVRRYNNNPNFSCALLKPEFSLI
 FADNHTELVAKTKHLDELQFKIMTGDIRYKSPAFSPLEIDAKCVGCPSCFESYSC
 NFQVINDTQVSGEGCTLFHNRI:ISANKSGYGIKMSCOTKPNONEFFIICNRRYS
 VLETT:DKNDK:EVNTGDOTSYIYKDSRCKTWCRCVREG"

BASE COUNT 225 a 358 c 349 g 279 t
 ORIGIN

Alignment Scores:
 Pred. No.: 9.45 Length: 1211
 Score: 43.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 88.89% Mismatches: 0
 Query Match: 97.73% Indels: 0
 DB: 14 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AY070271 (1-1211)

Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
 |||||
 Db 969 CTACTGGCGTCATGCTCAGCTC 995

RESULT 11
 AF187824/c
 LOCUS AF187824 2691 bp RNA linear VRL 02-MAY-2001
 DEFINITION Cache Valley virus 8068 GI gene, partial cds.
 ACCESSION AF187824
 VERSION AF187824.1 GI:6959495

KEYWORDS
 SOURCE
 ORGANISM
 Cache Valley virus
 Cache Valley virus
 Viruses; ssRNA negative-strand viruses; Bunyaviridae;
 Orthobunyavirus; Bunyamwera virus group.

REFERENCE
 1 (bases 1 to 2691)
 Brockus, C.L. and Grimstad, P.R.
 Comparative analysis of GI glycoprotein-coding sequences of Cache
 Valley virus (Bunyaviridae: Bunyavirus) isolates

JOURNAL
 MEDLINE 21221452
 PUBMED 11324749
 REFERENCE 2 (bases 1 to 2691)
 Brockus, C.L., Collins, F.H., Besansky, N.J., and Grimstad, P.R.
 Phylogenetic Analysis of Cache Valley Virus Isolates

TITLE
 JOURNAL
 REFERENCE 3 (bases 1 to 2691)
 Brockus, C.L. and Grimstad, P.R.
 Direct Submission
 TITLE
 JOURNAL
 Submitted (17-SEP-1999) Vector Biology Labs, University of Notre
 Dame, Galvin Life Sciences Center, Notre Dame, IN 46556, USA

FEATURES
 Location/Qualifiers
 1..2691
 /organism="Cache Valley virus"
 /mol_type="genomic RNA"

/strain="PC68"
 /db_xref="taxon:80935"
 /country="USA: Dennisville, NJ"
 /note="1982 isolate from Ae. sollicitans"
 1..2691
 /note="envelope glycoprotein; putative soluble portion"

GDS

/cddon_start=1
 /product="GI"
 /protein_id="AA533116.1"
 /db_xref="GI:6959496"

Alignment Scores:
 Pred. No.: 21.3 Length: 2691
 Score: 43.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 3
 Best Local Similarity: 88.89% Mismatches: 0
 Query Match: 97.73% Indels: 0
 DB: 14 Gaps: 0

BASE COUNT 949 a 476 c 539 g 727 t
 ORIGIN

US-09-965-594-1_COPY_14_22 (1-9) x AF187824 (1-2691)
 Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
 |||||
 Db 1367 TTGTTAGGGTGTGATACCTTCTCTG 1361

RESULT 12
 AF231116/c
 LOCUS AF231116 2691 bp DNA linear VRL 02-MAY-2001
 DEFINITION Cache Valley virus isolate Ar78-5912 GI envelope glycoprotein gene,
 partial cds.
 ACCESSION AF231116
 VERSION AF231116.1 GI:13249094

KEYWORDS
 SOURCE
 ORGANISM
 Cache Valley virus
 Cache Valley virus
 Viruses; ssRNA negative-strand viruses; Bunyaviridae;
 Orthobunyavirus; Bunyamwera virus group.

REFERENCE
 1 (bases 1 to 2691)
 Brockus, C.L. and Grimstad, P.R.
 Comparative analysis of GI glycoprotein-coding sequences of Cache
 Valley virus (Bunyaviridae: Bunyavirus) isolates

JOURNAL
 MEDLINE 21221452
 PUBMED 11324749
 REFERENCE 2 (bases 1 to 2691)
 Brockus, C.L. and Grimstad, P.R.
 Direct Submission
 TITLE
 JOURNAL
 Submitted (04-FEB-2000) Biological Sciences, University of Notre
 Dame, Galvin Life Sciences Bldg, Notre Dame, IN 46556, USA

FEATURES
 Location/Qualifiers
 1..2691
 /organism="Cache Valley virus"
 /mol_type="genomic DNA"
 /isolate="Ar78-5912"
 /specific_host="Anopheles quadrimaculatus"

```

/db_xref="taxon:50935"
/country="USA: Ohio, Kent"
CDS
<1..2691
/contig_start=1
/product="G1 envelope glycoprotein"
/protein_id="AAK15673.1"
/db_xref="GI:13249095"
/translation="ETDCWTKKSLDIDICGPMLDIGSCTNKNTSRDYDAEAKVVSQSK
ISQDAQVSLGNSIDSAIKAIRAOKTYPTMHLLETIFLMKNCXYTSFNHNSYSO
AKWRLAKTHGFDICSRHSHHFCRCINGCTQCONWDFAGEMNFTYNSKNDYFSD
LNFYIFENAFPTTESFYQIQKNTTGASKLGLKLTIKYGNKNMFGVWKFGGY
LFLPYINTELSQNRLLTATQELLSRSHSGROESMAIQGSITKFCNNAKGVGC
ISRFPGIPICNLTAGCSPNYKIVKPEKLYKNNKGEVWCNDVCLNDFEPADFEV
VKKIKLTCFLIDVSVVDVFSVAASCKMADKGVCTVNDKWNIIKCSGLYFTDH
REGQDNGDFGEYCISHSCNTRFPINPDIKCTWEPHRSKSKYSTISLESLEER
RAMAKUSHLLIYNKPTANPHIRPTTKFTANGVENSOGIESAYLSSIPALGGI
SVGVNQTQDNFPLDLIVFKSAIRTYTHIYDIGHTIGITQDHECTGCPAVV
PKENMTFSQERTSRMGCEFCCLATGCVFGSCDIHPETKYRKAVEEVLLI
VCINFPNNYCTEINAEIPKIDTEIQKFTVDTKLPNLLAVONHKLXSGOINDLGS
FSGCGNVQKTNHSLGMGTAKFDYVCHGASRKDVIVRCYNNKNSCKLLKEETSLI
FADNHETLEVAHKLHIGLGEQFKIMGLDIRYKFAESPELEIDAKCVGCSCEFSYSC
NFQIVNTDVTCSVEGPGCTLFPHNRILISANKQSYGLKMSCQTKPNONEEFFICNRKYS
VFITIDKNDKIEVNTGDTQSYIYEKDSRCKTWLCVRDEG"
BASE COUNT 947 a 475 c 541 g 728 t
ORIGIN
Alignment Scores:
Pred. No.: 21.3 Length: 2691
Score: 43.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 88.89% Mismatches: 0
Query Match: 97.73% Indels: 0
DB: 14 Gaps: 0
US-09-965-594-1_COPY_14_22 (1-3) x AF231118 (1-2691)
Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||
Db 1387 TTGTTAGGGTGTGTGATAACTTCCTC 1361

RESULT 13
AF231118/c
LOCUS Cache Valley virus isolate 61D240 G1 envelope glycoprotein gene,
DEFINITION partial cds.
ACCESSION AF231118
VERSION AF231118.1 GI:13249096
KEYWORDS
SOURCE Cache Valley virus
ORGANISM Cache Valley virus
VIRUSES: ssRNA negative-strand viruses; Bunyaviridae;
Orthobunyavirus; Bunyamwera virus group.
REFERENCE 1 (bases 1 to 2691)
AUTHORS Brockus,C.L. and Grimstad,P.R.
TITLE Comparative analysis of G1 glycoprotein-coding sequences of Cache
Valley virus (Bunyaviridae: Bunyavirus) isolates
JOURNAL Virus Genes 22 (2), 133-139 (2001)
MEDLINE 2122452
PUBMED 11324749
REFERENCE 2 (bases 1 to 2691)
AUTHORS Brockus,C.L. and Grimstad,P.R.
TITLE Phylogenetic analysis of Cache Valley virus isolates
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 4463)
AUTHORS Brockus,C.L. and Grimstad,P.R.
TITLE Direct Submission
JOURNAL Submitted (14-SEP-1999) Vector Biology Labs, University of Notre
Dame, Galvin Life Sciences Bldg, Notre Dame, IN 46556, USA
FEATURES
source
1..269;
/organism="Cache Valley virus"
/mol_type="genomic DNA"
/isolate="61D240"
/species="Bunyavirus"
/db_xref="taxon:50935"
/country="Mexico: Bocas el Toro"
/note="isolated in 1961

```


TCIC KASACAAKIQDCTKLVUGJDLVVICSAUTEDASIAVTEAKIKVSAPG
OPKPEDELEZSCSSNVAQSSAKRNVYLTPODTFLAKANATATAPVNSKL
GNIMVAPTLKAWIMLTHFESILLACQVLTQKALQVLTQVACYSILPELDTQLLHLC
GLSAPLSHVSPOETINVAALQRLGVLTILVNHKIRANVPKALQSSNHAANCQSYL
FNWAVRTKLTIPAAQSGJLSMFWACVSGGCVLTSLNAPNAPKPMCLLSLVY
GVLTIPNE"

BASE COUNT	1877 a	2830 c	2680 g	1968 f	4 others
ORIGIN					

Alignment Scores:	
Prod. No.:	76
Score:	43.02
Percent Similarity:	100.00%
Best Local Similarity:	88.89%
Query Match:	97.73%
DB:	14
Length:	1
Matches:	9
Conservative:	0
Mismatches:	1
Indels:	0
Gaps:	0
	9.559

US-09-965-594-1_COPY_14_22 (1-9) x AF313916 (2-9359)

Qy 1 leuLeuGlyCysIleIleThrSerLeu 9

3444 CTACTTGGTGCATCG:CAACAGGCTC 3470

Search completed: September 30, 2003, 12:51:59
Job time : 2207 secs

SanCore version 5.1.6
Copyright (c) 1993 - 2003 Computer Gen.

OM protein - nucleic search, using frame_plus_pin model

Run on: September 29, 2003, 19:09:03 : Search time 189 Seconds
(without alignments)
128,545 Million cell updates/sec

Title: us-09-965-594-1_copy_14_22

Perfect score: 44

Sequence: 1 L13C1:TSL 9

Scoring table:

R-OSUM62

Xgapop 10.0 , Xgapext 0.5

Ygapop 10.0 , Ygapext 0.5

Fgapop 6.0 , Fgapext 7.0

Delpop 6.0 , Delpext 7.0

Searched: 2552756 seqs, 1349719017 residues

Total number of hits satisfying chosen parameters: 5105271

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 99%

Listing first 45 summaries

Command line parameters:

-MODEL=frame+pin_model -DEV=xib

-O=Cg22_1/US10_spo01/US0995594/runat_29032003_154249_14068/app_query.fasta.1.199

-DB=N_Geneseq_19Jun03 -QMT=fastap -SUFFIX=ring -MINNA=CH-0.1 -LOOPEXT=0

-LIST=45 -DOCALIGN=200 -THR_SCORE=ptc -THR_MAX=99.9 -THR_MIN=0 -ALIGN=15

-MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000

-USER=US0995594 -CGN=1.1.312 -runat_29032003_104249_14068 -NCGU=6 -ICPU=3

-NO_MMAP -LARGEQUERY -NEG_SCORES=0 -WAIT -DSP512CK=100 -LONG3CG

-DEV_TIMEOUT=220 -WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FSGAPOP=6

-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

N_Geneseq_19Jun03:

1: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1980.DAT:

2: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1981.DAT:

3: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1982.DAT:

4: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1983.DAT:

5: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1984.DAT:

6: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1985.DAT:

7: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1986.DAT:

8: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1987.DAT:

9: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1988.DAT:

10: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1989.DAT:

11: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1990.DAT:

12: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1991.DAT:

13: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1992.DAT:

14: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1993.DAT:

15: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1994.DAT:

16: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1995.DAT:

17: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1996.DAT:

18: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1997.DAT:

19: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1998.DAT:

20: /SIDSL/gcdata/geneseq/geneseq-emb1/NA1999.DAT:

21: /SIDSL/gcdata/geneseq/geneseq-emb1/NA2000.DAT:

22: /SIDSL/gcdata/geneseq/geneseq-emb1/NA2001.DAT:

23: /SIDSL/gcdata/geneseq/geneseq-emb1/NA2002.DAT:

24: /SIDSL/gcdata/geneseq/geneseq-emb1/NA2003.DAT:

25: /SIDSL/gcdata/geneseq/geneseq-emb1/NA2003.DAT:

and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	43	97.7	9436	15	AA063499	Blood transmissible
2	42	95.5	819	15	AA058517	NANBH virus gene f
3	42	95.5	3481	15	AA064368	Non-A, non-B hepat
4	42	95.5	3481	15	AA050486	5'UTR/CORE/ENV/NS1
5	41	93.2	4977	23	AA086668	DNA encoding novel
6	41	93.2	4977	23	AA086891	DNA encoding novel
7	41	93.2	4977	23	AA089842	DNA encoding novel
8	41	93.2	9595	20	AA024843	Infectious hepatitis
9	41	93.2	9595	22	AA086939	Nucleotide sequenc
10	41	93.2	9595	22	AA023492	Infectious Hepatit
11	41	93.2	94895	21	AA022302	BAC containing rep
12	41	93.2	94895	21	AA022302	BAC containing rep
13	41	93.2	103929	21	AA022287	BAC containing rep
14	41	93.2	134499	21	AA022286	BAC containing rep
15	41	93.2	1082138	21	AA022305	Arabidopsis thalia
16	41	93.2	1082138	21	AA022305	Arabidopsis thalia
17	40	90.9	424	24	ABL63442	Breast cancer rela
18	40	90.9	1933	20	AA023258	HCV NS3 DNA. Hepa
19	40	90.9	2346	24	NA019166	cDNA encoding huma
20	40	90.9	2520	25	AB05429	Human novel polynu
21	40	90.9	8145	20	AA023259	Plasmid pET-BS(+)/
22	40	90.9	910715	20	AA020248	Porreilia burgdorfe
23	39	88.6	4154	25	AB019487	Aspergillus fumiga
24	39	88.6	4746	21	AA045995	Rat calcium/calmod
25	39	88.6	9402	14	AA041345	Human Hepatitis C
26	39	88.6	9405	14	AA040426	Full-length Hepati
27	38	86.4	242	25	AB023888	Human GDP-mannose
28	37	84.1	39	20	AA080326	HCV NS4A-NS3 compl
29	37	84.1	39	20	AA080327	HCV NS4A-NS3 compl
30	37	84.1	39	20	AA080328	HCV NS4A-NS3 compl
31	37	84.1	39	20	AA080329	HCV NS4A-NS3 compl
32	37	84.1	39	20	AA080330	HCV NS4A-NS3 compl
33	37	84.1	39	20	AA080331	HCV NS4A-NS3 compl
34	37	84.1	39	20	AA080332	HCV NS4A-NS3 compl
35	37	84.1	39	20	AA080333	HCV NS4A-NS3 compl
36	37	84.1	39	20	AA080334	HCV NS4A-NS3 compl
37	37	84.1	39	20	AA080335	HCV NS4A-NS3 compl
38	37	84.1	39	20	AA080316	HCV NS4A-NS3 compl
39	37	84.1	39	20	AA080317	HCV NS4A-NS3 compl
40	37	84.1	39	20	AA080318	HCV NS4A-NS3 compl
41	37	84.1	39	20	AA080319	HCV NS4A-NS3 compl
42	37	84.1	39	20	AA080320	HCV NS4A-NS3 compl
43	37	84.1	39	20	AA080321	HCV NS4A-NS3 compl
44	37	84.1	39	20	AA080322	HCV NS4A-NS3 compl
45	37	84.1	39	20	AA080323	HCV NS4A-NS3 compl

ALIGNMENTS

RESULT 1
AA063499

CD AA063499 standard; cDNA; 9435 BP.

XX AA063499;

AC AA063499;

BT 17-JAN-1995 (first entry)

XX Flood transmissible NANBHV genome.

DE Flood transmissible NANBHV genome.

XX Polymerase chain reaction; PCR; amplify; primer; non-A, non-B hepatitis;

KW NANBH; virus; blood transmissible; detection; hepatitis virus; RT-PCR;

KW C100 antibody; HCV RNA; NS5 region; ds.

XX Non-A, non-B hepatitis virus.

XX Key

PH Location/Qualifiers

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed.

FT CDS 342..9374
 FT /*tag= a

PN JP06105690-A.

PD 19-APR-1994.

XX 10-MAR-1992; 92JP-0051885.

XX 10-MAR-1992; 92JP-0051885.

XX (KAEN/) KAENNO K.

PA WPI; 1994-163139/20.

DR P-PSDB; AAR53417.

XX Blood-transmissible non-A non-B hepatitis virus DNA - used for
 PT detection of hepatitis virus

PS Claim 1; Page 8-20; 22pp; Japanese.

XX This sequence represents the genome of a blood transmissible non-A.
 CC non-B hepatitis (NANBH) virus. This sequence was isolated using the
 CC primers given in AAQ63500-95. The amplified fragments are used in the
 CC detection of hepatitis virus. This target DNA was isolated from serum
 CC of chronically infected NANBH patients who were C100 antibody-positive
 CC and HCV RNA (NS5 region) positive. Reverse transcription-PCR and PCR
 CC were performed on cDNA and the total human NANBH DNA was constructed
 CC from 23 clones.

XX Sequence 9436 BP; 1876 A; 2840 C; 2055 G; 1974 T; 91 other;

Alignment Scores:
 Pred. No.: 653 Length: 9436
 Score: 43.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 1
 Best Local Similarity: 88.89% Mismatches: 0
 Query Match: 97.73% Indels: 0
 DB: 15 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAQ63459 (1-9436)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9

DB 3456 CTAAGTGGTGCATTGTACTAGCCTC 3482

RESULT 2

AAQ58817

ID AAQ58817 standard; cDNA: 819 BP.

XX AC AAQ58817;

DI 24-NOV-1994 (first entry)

DE NANBH virus gene fragment #4.

XX Antigen; structural; non-structural; non A non B hepatitis virus;
 KW NANBH; NANBH; patient; plasma; diagnosis; detection; carrier; ss.

XX Non A Non B hepatitis virus.

PN JP06070778-A.

XX 15-MAR-1994.

PF 01-JUN-1993; 92JP-0156087.

XX 10-JUL-1992; 92JP-0207391.

XX (KOKU-) KOKUSAI SHIYAKU KK.

PA (SANW) SANWA KAGAKU KENKYUSHO CO.

PA (TOFU) TONEN CORP.

PA (TOKR-) 2H TOKYOOTO RINSHO IGAKU SOGO KENKYUSHO.

XX WPI; 1994-128677/16.
 DR P-PSDB; AAR50072.

XX Nucleic acid fragment coding non-A non-B hepatitis virus antigen
 PT - useful in diagnosis of NANB patient and detection of virus
 PT carrier

XX Claim 8; Page 18-19; 37pp; Japanese.

XX The sequences given in AAQ58814-27 encode antigens of structural and
 CC non-structural regions of non A non B hepatitis virus (NANBHV).
 CC These sequences were derived from the plasma of a NANBH patient by
 CC recombinant DNA techniques. These fragments are useful for the
 CC diagnosis of NANBH patients and the detection of NANBH carriers.

XX Sequence 819 BP; 150 A; 255 C; 241 G; 173 T; 0 other;

Alignment Scores:
 Pred. No.: 66.4 Length: 819
 Score: 42.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 1
 Best Local Similarity: 88.89% Mismatches: 0
 Query Match: 95.45% Indels: 0
 DB: 15 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAQ58817 (1-819)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9

DB 67 CTAATGGCTGCATCATCAGCAGCCTC 93

RESULT 1

AAQ64068

ID AAQ64068 standard; cDNA: 3461 BP.

XX AC AAQ64068;

DI 14-FEB-1995 (first entry)

DE Non-A, non-B hepatitis virus gene #4.

XX Non-A, non-B hepatitis virus; NANBHV; hepatitis C virus; HCV;
 KW core; ENV; NS1; NS2; NS3; antigen; detection; ss.

XX Hepatitis C virus.

XX Key

FT CDS Location/Qualifiers

FT 307..3461

FT /*tag= a

FT misc_RNA 307..879

FT /*tag= b

FT misc_RNA 880..1455

FT /*tag= c

FT misc_RNA 1456..2736

FT /*tag= d

FT misc_RNA 2737..3461

FT /*tag= e

FT /*tag= NS2-3

FT /*tag= "NS3 N-terminal"

PN JP06141870-A.

XX 24-MAY-1994.

PF 12-MAR-1992; 92JP-0088140.

XX 12-MAR-1992; 92JP-0088140.

XX (SANW) SANWA KAGAKU KENKYUSHO CO.

PA (TOFU) TONEN CORP.
 PA (TOKR) 2H TOKYOTO RINSHO IGAKU SOGOKU KENKYUSEI.
 XX
 DR WPI: 1994-225026/25.
 DR P-PSDB: AAR54066.
 XX
 XX DNA coding a Non-A, non-B hepatitis virus antigen - useful for
 PT detecting HCV within serum
 XX
 PS Claim 1-5; Page 11-15; 22pp; Japanese.
 XX
 CC Hepatitis C virus #4 and #6 genes were isolated (AA064068-69).
 CC Both genes contain the core, ENV, NS1, NS2 and NS3 regions.
 CC A core region fragment is given in AA064067.
 XX
 SQ Sequence 3461 BP; 638 A; 1046 C; 1012 G; 765 T; 0 other;

Alignment Scores:
 Pred. No.: 336 Length: 3461
 Score: 42.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 1
 Best Local Similarity: 88.99% Mismatches: 0
 Query Match: 95.45% Indels: 0
 DB: 15 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AA064068 (1-3461)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
 |||||:|||||:|||||:|||||:|||||

DB 3421 CTAATTGGCTGCATCATCACCAGCCTA 3447

RESULT 4
 AA030386
 ID AA030386 standard: cDNA: 3461 BP.
 XX
 AC AA030386;
 XX
 DT 22-AUG-1996 (first entry)
 XX
 DE 5'UTR/CORE/ENV/NS1/NS2/NS3 cDNA from HCV (#4).
 XX
 KW Hepatitis C virus; HCV; antigen; detection; antibody; dis.
 XX
 OS Hepatitis C virus.
 XX
 FH Key Location/Qualifiers
 FT CDS 307..3461
 FT /*tag= a
 FT misc_feature 307..373
 FT /*tag= b
 FT /*product= core peptide
 FT misc_feature 880..1455
 FT /*tag= c
 FT /*product= ENV;
 FT misc_feature 1456..2736
 FT /*tag= d
 FT /*product= NS1/ENV2
 FT misc_feature 2737..3461
 FT /*tag= e
 FT /*product= NS2 and NS3

JP07133291-A.
 23-MAY-1995.
 18-JUN-1993; 93JP-0147944.
 16-JUN-1993; 93JP-0147944.
 XX (TOFU) TONEN CORP.
 PA
 XX WPI: 1995-220780/29.
 DR P-PSDB: AAR98361.

XX Recombinant polypeptide comprising partial NS1 region of hepatitis
 PT non-A non-B viral antigen - used in a method for detecting
 PT antibodies against hepatitis non-A non-B virus.
 XX
 PS Disclosure: Page 10-12; 15pp; Japanese.
 XX
 CC The sequences given in AA030386-87 encode the 5'UTR/CORE/ENV/NS1/NS2/
 CC NS3 protein region derived from hepatitis C virus (HCV) isolates #4/
 CC and #6 respectively. The proteins encoded by these sequences partic.
 CC encode amino acids 394-495 of the HCV NS1 antigen. These protein
 CC fragments may be used in the detection of antibodies against HCV.
 XX
 SQ Sequence 3461 BP; 638 A; 1046 C; 1012 G; 765 T; 0 other;

Alignment Scores:
 Pred. No.: 336 Length: 3461
 Score: 42.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 1
 Best Local Similarity: 88.89% Mismatches: 0
 Query Match: 95.45% Indels: 0
 DB: 16 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AA030386 (1-3461)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
 |||||:|||||:|||||:|||||:|||||

DB 3421 CTAATTGGCTGCATCATCACCAGCCTA 3447

RESULT 5
 AA086668/C
 ID AA086668 standard: cDNA: 4977 BP.
 XX
 AC AA086668;
 XX
 DT 13-FEB-2002 (first entry)
 XX
 DE DNA encoding novel human diagnostic protein #22472.
 XX
 KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX
 CS Homo sapiens.
 XX
 PN NC0200175067-A2.
 XX
 PC 11-DEC-2001.
 XX
 PF 30-MAR-2001; 2001WG-US08631.
 XX
 PR 31-MAR-2000; 2000US-0540217.
 PR 23-AUG-2000; 2000US-0649167.
 XX
 PA (HYSEP-) HYSEQ INC.
 XX
 PI Dimasac RT, Liu C, Tang YT;
 XX
 DR WPI: 2001-639362/73.
 DR P-PSDB: AB022481.
 XX
 XX Now isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 XX
 PS Claim 1: SEQ ID NO 22472; 103pp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques

CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 4977 BP: 1417 A: 1392 C: 1202 G: 966 T: 0 other:

Alignment Scores:
 Pred. No.: 795 Length: 4977
 Score: 41.00 Matches: 7
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 77.78% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 23 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAS86658 (1-4977)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9

DB 1530 CTTTGGGCGTGTTCCTTACTTCCTG 1504

RESULT: 6

AAS88891/C

ID AAS88891 standard; cDNA: 4977 BP.

XX AAS88891;

DI 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #24595.

XX Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.

XX Homo sapiens.

XX WO200175067-A2.

XX 11-OCT-2001.

XX 30-MAR-2001; 2001WO-US08631.

XX 31-MAR-2000; 2000US-0540217.

XX 23-AUG-2000; 2000US-0649167.

XX (HYSE-) HYSEQ INC.

XX Drmanac RT, Liu C, Tang YT;

XX WPI: 2001-639362/73.

XX P-PSDB; ABG24704.

XX New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.

XX Claim 1: SEQ ID No 24595; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and

CC polypeptide (II) sequences. (I) is useful as hybridisation probes,

CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome

CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC the polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 4977 BP: 1417 A: 1392 C: 1202 G: 966 T: 0 other:

Alignment Scores:
 Pred. No.: 795 Length: 4977
 Score: 41.00 Matches: 7
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 77.78% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 23 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAS88891 (1-4977)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9

DB 1530 CTTTGGGCGTGTTCCTTACTTCCTG 1504

RESULT: 7

AAS88891/C

ID AAS88891 standard; cDNA: 4977 BP.

XX AAS88891;

DI 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #25646.

XX Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.

XX Homo sapiens.

XX WO200175067-A2.

XX 11-OCT-2001.

XX 30-MAR-2001; 2001WO-US08631.

XX 31-MAR-2000; 2000US-0540217.

XX 23-AUG-2000; 2000US-0649167.

XX (HYSE-) HYSEQ INC.

XX Drmanac RT, Liu C, Tang YT;

XX WPI: 2001-639362/73.

XX P-PSDB; ABG25655.

XX New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.

XX Claim 1: SEQ ID No 25646; 103pp; English.

CC The invention relates to isolated polypeptide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes.
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (I). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against its detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (I) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS8457-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from W:PO
 CC at ftp.wipo.int/pub/published_pat_sequences.

XX Sequence 4977 BP; 1417 A; 1392 C; 1202 G; 966 T; 0 other;

Alignment Scores:
 Pred. No.: 795 Length: 4977
 Score: 41.00 Matches: 7
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 77.78% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 23 Gaps: 0

US-09-965-594-1_copy_14_22 (1-9) x AAS89842 (1-4977)

Cy 1 LeuLeuGlyCysIleIleThrSerLeu 9

Db 1530 CTTTGGGAGTGATTCAGCTCTCTG 1504

RESULT 9

AAX24843

ID AAX24843 standard; DNA: 9595 BP.

XX AAX24843;

XX 21-JUN-1999 (first entry)

DE Infectious hepatitis C virus genotype 1b strain HC-J4 genome.

XX HCV; infectious clone; infection; diagnosis; therapy; vaccine;
 KW screening; assay; antiviral; virucide; ss.

XX Hepatitis C virus.

FE Key Location/Qualifiers
 FT CDS 342..9374
 FT /*tag- a

PN WO9901008-A2.

PD 28-JAN-1999.

XX 16-JUL-1998; 96WO-US14588.

PR 27-JAN-1996; 98JUS-C034416.

PR 18-JUL-1997; 97US-C053062.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Buxh J, Emerson SU, Purcell RH, Yanagi M;

XX WPI; 1999-132252/11.

XX P-PSDB; AAW95022.

XX

PT New isolated hepatitis C virus nucleic acids - used to develop
 PT products for the diagnosis, prevention and treatment of HCV
 PT infections and for developing screening assays

PS claim 3; Fig 1A-E; 126pp; English.

XX The present sequence comprises the nucleic acid sequence of the
 CC genome of infectious hepatitis C virus (HCV) genotype 1b strain
 CC HC-J4 (ATCC 29596); that is capable of expressing this virus when
 CC transfected into cells. HC-J4 was obtained from acute phase plasma
 CC of a chimpanzee experimentally infected with serum containing
 CC HC-J4/91. The claimed infectious nucleic acid sequence can be used
 CC to produce chimeric genomes (see AAX24843) consisting of the open
 CC reading frames of infectious nucleic acid sequences of other
 CC genotypes (including genotypes 1-6) and subtypes (such as 1b, 2a,
 CC 2b, 2c, 3a, 4a-f, 5a and 6a) of HCV. The invention also relates to
 CC the introduction of mutations or deletions into infectious nucleic
 CC acid sequences in order to produce an attenuated HCV virus suitable
 CC for vaccine development. Infectious nucleic acid sequences can
 CC also be used to produce attenuated virus via passage in vitro or in
 CC vivo of the viruses produced by transfection of a host cell with
 CC the infectious nucleic acid sequence. Vaccines comprising one or
 CC more polypeptides made from the infectious nucleic acid sequence are
 CC used to immunise mammals, especially humans, against hepatitis C.
 CC The nucleic acid sequences can also be used to induce protective
 CC immunity against the virus. The nucleic acid sequences or their
 CC encoded proteases (e.g. NS3 protease) can additionally be used to
 CC develop screening assays to identify antiviral agents for HCV.

XX Sequence 9595 BP; 1934 A; 2842 C; 2698 G; 2121 T; 0 other;

Alignment Scores:

Pred. No.: 1-66e+03 Length: 9595
 Score: 41.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 1
 Best Local Similarity: 88.89% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 20 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAX24843 (1-9595)

Cy 1 LeuLeuGlyCysIleIleThrSerLeu 9

Db 3456 GTACITGGTTCATCATCACTAGCCCTC 3482

RESULT 9

AAC86939

ID AAC86939 standard; DNA: 9595 BP.

XX AAC86939;

XX 02-APR-2001 (first entry)

XX Nucleotide sequence of a hepatitis C virus (HCV) clone genotype 1b.

XX Chimeric virus; bovine viral diarrhoea virus; BVDV; hepatitis C virus;
 KW HCV; vaccine; viral inhibitor; antiviral; ss.

XX Hepatitis C virus.

FE Key Location/Qualifiers
 FT CDS 342..9374
 FT /*tag- a

PN WO200075352-A2.

XX 14-DEC-2000.

XX 02-JUN-2000; 2000WO-US15527.

XX 04-JUN-1999; 99US-0137917.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Nam J, Bukh J, Emerson SJ, Purcell RH;
 PI WPI: 2001-071061/C8.
 DR P-PSDB; AAB31170.

XX New nucleic acid comprising a chimeric bovine viral diarrhoea virus
 PT genome in which the (non-)structural region has been replaced by
 PT hepatitis C virus (HCV) genome useful for treating or preventing HCV
 PT signs and symptoms

XX Disclosure; Fig 4A-F; 97pp; English.

XX The specification describes a nucleic acid comprising a chimeric virus
 CC genome, specifically bovine viral diarrhoea virus (BVDV) genome in which
 CC the (non-)structural region has been replaced by the (non-)structural
 CC region of a hepatitis C virus (HCV) genome. The nucleic acids comprising
 CC the chimeric virus and the chimeric virus are useful for identifying
 CC cell lines capable of supporting the replication of these chimeric
 CC viruses, in screening for neutralizing antibodies to HCV of different
 CC genotypes, in the production of HCV-BVDV viruses, for the development
 CC of inactivated or attenuated vaccines to prevent HCV-BVDV in a mammal,
 CC in studying the molecular properties of HCV indirectly in vitro, and in
 CC identifying inhibitors of viral enzyme activity which would be useful
 CC as antiviral agents. Formulations of compositions comprising the
 CC chimeric viruses may be used to treat or prevent the signs and symptoms
 CC of HCV. The present sequence represents a BVDV clone, which is used
 CC to construct chimeric nucleic acids of the invention.

XX Sequence 9595 BP; 1934 A; 2843 C; 2697 G; 2121 T; 0 other;

Alignment Scores:
 Pred. No.: 1,660+03 Length: 9595
 Score: 41.00 Matches: 8
 Percent Similarity: 100.00% Conservatve: 1
 Best Local Similarity: 88.89% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 22 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAC66939 (1-9595)

Oy 1 LeuLeuGlyCysIleIleThrSerLeu 9

Db 3456 GTACTTGTTGCATCATCTAGCCGTC 3482

RESULT 10

AAP23492

ID AAP23492 standard; DNA; 9595 BP.

AC AAP23492;

XX 21-MAR-2001 (first entry)

XX Infectious Hepatitis C virus 1b genotype.

DE GBV-B; hepatitis C virus; HCV; vaccine; ds.

XX Hepatitis C virus.

OS WO200075337-A1.

PN 14-DEC-2000.

XX 02-JUN-2000; 2000WO-US15293.

XX 04-JUN-1999; 99US-0137694.

XX (US) US DEPT HEALTH & HUMAN SERVICES.

XX Bukh J, Yanagi M, Emerson SJ, Purcell RH;

XX WPI: 2001-091214/10.

XX

PT New infectious nucleic acids of the GB virus-B clone, useful for
 PT indirectly studying the molecular properties of hepatitis C virus (HCV)
 PT and in developing vaccines and therapeutics for HCV
 XX Disclosure; Fig 7; 96pp; English.

XX The present invention relates to GB virus-B. The nucleic acid molecules
 CC of the invention are useful for indirectly studying the molecular
 CC properties of hepatitis C virus (HCV). The infectious nucleic acid
 CC sequence of the GB virus-B clone and the HCV/GBV-B chimeras may be used
 CC in the development of vaccines and therapeutics for HCV.

XX Sequence 9595 BP; 1934 A; 2843 C; 2697 G; 2121 T; 0 other;

Alignment Scores:
 Pred. No.: 1,660+03 Length: 9595
 Score: 41.00 Matches: 8
 Percent Similarity: 100.00% Conservatve: 1
 Best Local Similarity: 88.89% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 22 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAP23492 (1-9595)

Oy 1 LeuLeuGlyCysIleIleThrSerLeu 9

Db 3456 GTACTTGTTGCATCATCTAGCCGTC 3482

RESULT 1:

AAP23492

ID AAP23492 standard; DNA; 94895 BP.

AC AAP23492;

XX 20-MAR-2001 (first entry)

XX BAC containing repeats from centromeres 1-4 #25.

XX Centromere; microsome; vector; ds.

OS Arabidopsis thaliana.

XX WO200055325-A2.

XX 21-SEP-2000.

XX 17-MAR-2000; 2000WO-US07392.

XX 18-MAR-1999; 99US-0125219.

XX 01-APR-1999; 99US-0127409.

XX 18-MAY-1999; 99US-0134770.

XX 13-SEP-1999; 99US-0153584.

XX 17-SEP-1999; 99US-0154603.

XX (UYCH-) UNIV CHICAGO.

XX Preuss D, Copenhaver G, Keith K;

XX WPI: 2000-587529/55.

XX Recombinant DNA construct comprising a plant centromere, useful for
 PT producing stably inherited microsome which can serve as vectors for
 PT the construction of transgenic plant and animal cells -
 XX Claim 102; Page 804-819; 1449pp; English.

XX The present invention relates to a recombinant DNA construct of a plant
 CC (Arabidopsis thaliana) centromere. The constructs are useful for
 CC producing stably inherited microsome which can serve as vectors for
 CC the construction of transgenic plant and animal cells expressing
 CC selected proteins such as hormones, enzymes, interleukins, clotting
 CC factors, cytokines, antibodies, and growth factors.

US-09-965-594-1_COPY_14_22 (1-5) x AAF22302 (1-94895)

DE BAC containing repeats from centromeres 1-4 #9.
 XX
 KW Centromere: microsome; vector; ds.
 XX
 XX Arabidopsis thaliana.
 OS
 PN WO200055325-A2.
 XX
 XX 21-SEP-2000.
 XX
 XX 17-MAR-2000; 2000WO-US07392.
 PF
 XX 18-MAR-1999; 99US-0125219.
 PR
 PR 01-APR-1999; 99US-0127459.
 PR
 PR 18-MAY-1999; 99US-0134770.
 PR
 PR 13-SEP-1999; 99US-0154584.
 PR
 PR 17-SEP-1999; 99US-0154603.
 XX
 XX (UYCH-) UNIV CHICAGO.
 PA
 XX Preuss D, Copenhaver G, Keith K;
 PI
 XX WPI: 2000-587529/55.
 DR
 XX Recombinant DNA construct comprising a plant centromere, useful for
 PT producing stably inherited microsomies which can serve as vectors for
 PI the construction of transgenic plant and animal cells
 DR
 XX Claim 102; Page 451-484; 1449pp; English.
 PS
 XX The present invention relates to a recombinant DNA construct of a plant
 CC (Arabidopsis thaliana) centromere. The constructs are useful for
 CC producing stably inherited microsomies which can serve as vectors for
 CC the construction of transgenic plant and animal cells expressing
 CC selected proteins such as hormones, enzymes, interleukins, clotting
 CC factors, cytokines, antibodies, and growth factors.
 CC
 XX Sequence 134499 BP; 41565 A; 25130 C; 25225 G; 42577 T; 2 other;
 SQ
 Alignment Scores:
 Pred. No.: 3-25e-04 Length: 134499
 Score: 41.00 Matches: 7
 Percent Similarity: 100.00% Conservatives: 2
 Best Local Similarity: 77.78% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 21 Gaps: 0
 US-09-965-594-l_COPY_14_22 (1-9) x AAF22286 (1-134499)
 QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
 DB 1360 TTGTTAGGTTCATTGTCGACGAGTATG 1386
 RESULT 15
 AAF22305
 ID AAF22305 standard; DNA; 1082138 BP.
 AC
 XX AAF22305;
 XX
 XX 20-MAR-2001 (first entry)
 DT
 XX Arabidopsis thaliana chromosome 4 centromere.
 DE
 XX Centromere: microsome; vector; ds.
 XX
 XX Arabidopsis thaliana.
 OS
 PN WO200055325-A2.
 XX
 XX 21-SEP-2000.
 XX
 XX 17-MAR-2000; 2000WO-US07392.
 PF
 XX 18-MAR-1999; 99US-0125219.
 PR
 PR 01-APR-1999; 99US-0127459.
 PR
 PR 18-MAY-1999; 99US-0134770.
 PR
 PR 13-SEP-1999; 99US-0154584.
 PR
 PR 17-SEP-1999; 99US-0154603.
 XX
 XX (UYCH-) UNIV CHICAGO.
 PA
 XX Preuss D, Copenhaver G, Keith K;
 PI
 XX WPI: 2000-587529/55.
 DR
 XX Recombinant DNA construct comprising a plant centromere, useful for
 PT producing stably inherited microsomies which can serve as vectors for
 PI the construction of transgenic plant and animal cells
 DR
 XX Claim 102; Page 451-484; 1449pp; English.
 PS
 XX The present invention relates to a recombinant DNA construct of a plant
 CC (Arabidopsis thaliana) centromere. The constructs are useful for
 CC producing stably inherited microsomies which can serve as vectors for
 CC the construction of transgenic plant and animal cells expressing
 CC selected proteins such as hormones, enzymes, interleukins, clotting
 CC factors, cytokines, antibodies, and growth factors.
 CC
 XX Sequence 134499 BP; 41565 A; 25130 C; 25225 G; 42577 T; 2 other;
 SQ

PR 18-MAR-1999; 99US-0125219.
 PR
 PR 01-APR-1999; 99US-0127459.
 PR
 PR 18-MAY-1999; 99US-0134770.
 PR
 PR 13-SEP-1999; 99US-0154584.
 PR
 PR 17-SEP-1999; 99US-0154603.
 XX
 XX (UYCH-) UNIV CHICAGO.
 PA
 XX Preuss D, Copenhaver G, Keith K;
 PI
 XX WPI: 2000-587529/55.
 DR
 XX Recombinant DNA construct comprising a plant centromere, useful for
 PT producing stably inherited microsomies which can serve as vectors for
 PI the construction of transgenic plant and animal cells
 DR
 XX Claim 68; Page 977-1388; 1449pp; English.
 PS
 XX The present invention relates to a recombinant DNA construct of a plant
 CC (Arabidopsis thaliana) centromere. The constructs are useful for
 CC producing stably inherited microsomies which can serve as vectors for
 CC the construction of transgenic plant and animal cells expressing
 CC selected proteins such as hormones, enzymes, interleukins, clotting
 CC factors, cytokines, antibodies, and growth factors.
 CC
 XX Sequence 1082138 BP; 348775 A; 194404 C; 195515 G; 343444 T; 0 other;
 SQ
 Alignment Scores:
 Pred. No.: 3-29e+05 Length: 1082138
 Score: 41.00 Matches: 7
 Percent Similarity: 100.00% Conservatives: 2
 Best Local Similarity: 77.78% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 21 Gaps: 0
 US-09-965-594-l_COPY_14_22 (1-9) x AAF22305 (1-1082138)
 QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
 DB 885359 TTGTAGCTTCATTGTCGACGAGTATG 885065
 Search completed: September 29, 2003, 19:19:33
 Job time : 244 secs

GenCore version 5.1.1
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - nucleic search, using frame_plus_p2a.cole1

Run on: September 29, 2003, 19:11:08 : Search time 1425 seconds
(without alignments)
134,539 Million cell updates/sec

Title: US-09-965-594-1_COPY_14_22

Perfect score: 44

Sequence: 1 LOCIIIT5.9

Scoring table: BLOSUM62

Xgapop 10.0, Xgapext 0.5

Ygapop 10.0, Ygapext 0.5

Fgapop 6.0, Fgapext 7.0

Delop 5.0, Delext 7.0

Searched: 22781392 seqs, 12152238056 residues

Total number of hits satisfying chosen parameters: 45562778

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 99%

Listing first 45 summaries

Command line parameters:

-MODEL-frame+ p2n.model -DEV-x1x

-Q/cgn2_1/USFPO-spool/US09965594/runat_29092003_164249_14082/app_query.fasta_1.159

-DB-EST -QFMT=fastap -SUFFIX=rst -MINMATCH=0.1 -LOOPCL=0 -LOOPEXT=0

-UNITS-bits -SPART=1 -MATRIX=blosum62 -TRANS-human40.cdi -LIST=45

-DOCALIGN=200 -THR_SCORE=pct -THR_MAX=99.9 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL

-OUTFMT=ptc -NORM-ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000

-USER-US09965594 -ACGN_1_1_2810 -runat_29092003_164249_14082 -NUP=6 -ICPU=3

-NO_MMAL -LARGEQUERY -NEG_SCORES=0 -WAIT -DSPBLOCK=100 -LONLOG

-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6

-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

1: em_estab1.*

2: cm_estab1.*

3: cm_estab1.*

4: cm_estab1.*

5: em_estab1.*

6: em_estab1.*

7: em_estab1.*

8: cm_estab1.*

9: qb_estab1.*

10: qb_estab1.*

11: qb_estab1.*

12: qb_estab1.*

13: qb_estab1.*

14: qb_estab1.*

15: cm_estab1.*

16: em_estab1.*

17: cm_estab1.*

18: cm_estab1.*

19: em_estab1.*

20: em_estab1.*

21: em_estab1.*

22: em_estab1.*

23: em_estab1.*

24: cm_estab1.*

25: em_estab1.*

26: cm_estab1.*

27: em_estab1.*

28: qb_estab1.*

29: gb_gss2.*

Prod. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	% Match	Query Length	DB ID	Description
C 1	41	93.2	108	28	BH241427
C 2	41	93.2	240	29	AL950918
C 3	41	93.2	334	29	BZ352875
C 4	41	93.2	400	29	CNS00021
C 5	41	93.2	424	28	BH758372
C 6	41	93.2	444	10	BF651538
C 7	41	93.2	465	10	BF651537
C 8	41	93.2	498	28	B96162
C 9	41	93.2	505	28	B26036
C 10	41	93.2	631	29	BZ289143
C 11	41	93.2	683	28	BH241556
C 12	41	93.2	693	28	B95974
C 13	41	93.2	719	28	BH241494
C 14	41	93.2	724	28	AO955329
C 15	41	93.2	758	28	BH241165
C 16	41	93.2	787	28	BH241117
C 17	41	93.2	823	28	BH241591
C 18	41	93.2	831	28	BH241228
C 19	41	93.2	835	29	CC429018
C 20	41	93.2	847	29	CC363288
C 21	41	93.2	917	10	BF167496
C 22	40	90.9	287	9	AI800243
C 23	40	90.9	309	9	AI587235
C 24	40	90.9	355	14	Z39665
C 25	40	90.9	384	9	AI583735
C 26	40	90.9	401	14	CA902724
C 27	40	90.9	406	10	BF109646
C 28	40	90.9	407	10	BF727441
C 29	40	90.9	413	28	AQ759141
C 30	40	90.9	423	9	AA721993
C 31	40	90.9	424	14	TC3709
C 32	40	90.9	425	9	AA812004
C 33	40	90.9	426	9	AA665172
C 34	40	90.9	473	9	AI927322
C 35	40	90.9	540	12	BQ017316
C 36	40	90.9	588	10	BE785099
C 37	40	90.9	588	14	HI5452
C 38	40	90.9	613	14	HI5273
C 39	40	90.9	625	12	BM682281
C 40	40	90.9	683	29	BZ325209
C 41	40	90.9	702	13	BQ184273
C 42	40	90.9	710	13	BQ187309
C 43	40	90.9	915	10	BG685367
C 44	40	90.9	927	13	BU527696
C 45	40	90.9	1006	10	BG758345

ALIGNMENTS

RESULT 1	108 bp	DNA	linear	GSS 13-NOV-2001
BH241427/c				
LOCUS				
DEFINITION	BH241427	Arabis thaliana genomic clone	AU1LF555	genomic
ACCESSION	BH241427	survey sequence		
VERSION	BH241427.1	GI:16915855		
KEYWORDS	GSS			
SOURCE	Arabis thaliana (thale cress)			
ORGANISM	Arabis thaliana			
	Eukaryota; Viridiplantae;			
	Spermatophyta; Magnoliophyta; core eudicots; rosids			
	; eurosid II; Brassicales; Brassicaceae; Arabidopsi			

REFERENCE 1 (bases 1 to 106);
 AUTHORS Town.C.D., Whitelaw,C.A., Pai,G., Van Aken,S.E., Utterback,T.V.,
 Feldblyum,T.V. and Fraser,C.M.
 TITLE Survey sequencing of Arabidopsis thaliana BAC F26J21
 JOURNAL Unpublished
 COMMENT Other_GSSs: AUILF55TF
 Contact: Chris Town
 tIGR
 3712 Medical Center Drive, Rockville, MD 20850, USA.
 Tel: 301-838-3523
 Fax: 301-838-0208
 Email: cdtown@tigr.org
 From Wash. U contig 720.
 Seq primer: TR
 Class: sheared ends.

FEATURES
 source
 1..108
 Location/Qualifiers
 /organism="Arabidopsis thaliana"
 /mol_type="genomic DNA"
 /strain="Columbia 0"
 /db_xref="taxon:3702"
 /clone="AUILF55"
 /clone_lib="AUIL"
 /note="Vector: pHC82; Site_1: 2-3 kb sheared BAC
 DNA inserted into pHC82 using BstXI linkers"
 BASE COUNT 36 a 29 c 8 g 35 t
 ORIGIN
 Alignment Scores:
 Pred. No.: 90.9 Length: 106
 Score: 41.00 Matches: 7
 Percent Similarity: 100.00% Conservat.: 2
 Best Local Similarity: 77.78% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 28 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x B241427 (1-103)
 QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
 |||||
 DB 56 TTGTAGGTCATGTCGACGAGTATG 39

RESULT 2
 AL950918
 LOCUS Arabidopsis thaliana 230 bp DNA linear GSS 24-OCT-2002
 DEFINITION Arabidopsis thaliana T-DNA flanking sequence GK-332C10-016064,
 genomic survey sequence.
 ACCESSION AL950918.1 GI:24407540
 VERSION A:950918
 KEYWORDS GSS.
 SOURCE Arabidopsis thaliana (thale cress)
 ORGANISM Arabidopsis thaliana
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 rosids; eustosids II; Brassicales; Brassicaceae; Arabidopsis.
 REFERENCE 1
 AUTHORS Strizhov,N., Li,Y., Rosso,M., Viehoever,P., Dekker,K., Saedler,H.
 and Weissshaar,B.
 TITLE A pipeline for automated high-throughput generation of FSTs
 (flanking sequence tags) from Arabidopsis thaliana T-DNA
 transformed lines
 JOURNAL Unpublished
 REFERENCE 2
 AUTHORS Rosso,M., Strizhov,N., Li,Y., Reiss,B., Dekker,K. and Weissshaar,B.
 TITLE A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat)
 for flanking sequence tag based reverse genetics
 JOURNAL Unpublished
 REFERENCE 3 (bases 1 to 230)
 AUTHORS Li,Y., Rosso,M., Strizhov,N. and Weissshaar,B.
 TITLE Direct Submission
 JOURNAL Submitted (21-OCT-2002) Weissshaar B., Max-Planck-Institut fuer
 Zuechtungsforschung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany
 COMMENT This sequence is recovered from the left border of the T-DNA. It

indicates an insertion within the locus defined by clone FAmly. The
 sequences are generated at the MPI for Plant Breeding Research in
 the context of the GABI-Kat project. GABI-Kat is part of the German
 Plant Genomics program designated "GABI". Information on line
 availability can be found at:
 http://www.mpiz-koeln.mpg.de/GABI-Kat/.

FEATURES
 source
 1..230
 Location/Qualifiers
 /organism="Arabidopsis thaliana"
 /mol_type="genomic DNA"
 /strain="Columbia 0"
 /db_xref="taxon:3702"
 /clone="GK-332C10-016064"
 /clone_lib="Arabidopsis thaliana T-DNA insertion lines"
 /note="PCR was performed on DNA from Arabidopsis thaliana
 plants (T1) which were transformed with the T-DNA from
 vector PAC161. The lines contain one or more T-DNA
 insertions. The DNA fragment(s) resulting from the PCR
 were directly sequenced to determine the genomic sequence
 flanking the insertion. Sequences displaying significant
 similarity to the A. thaliana nuclear genome sequence were
 processed for submission. T-DNA derived sequences were
 removed"
 BASE COUNT 67 a 42 c 48 g 73 t
 ORIGIN
 Alignment Scores:
 Pred. No.: 277 Length: 230
 Score: 41.00 Matches: 7
 Percent Similarity: 100.00% Conservat.: 2
 Best Local Similarity: 77.78% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 29 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AL950918 (1-230)
 QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
 |||||
 DB 164 TTGTAGGTCATGTCGACGAGTATG 190

RESULT 3
 B2352875/C
 LOCUS Arabidopsis thaliana 334 bp DNA linear GSS 14-NOV-2002
 DEFINITION SALK_082734.11.45.x Arabidopsis thaliana T-DNA insertion lines
 Arabidopsis thaliana genomic clone SALK_082734.31.45.x, genomic
 survey sequence.
 ACCESSION B2352875.1 GI:24943737
 VERSION B2352875
 KEYWORDS GSS.
 SOURCE Arabidopsis thaliana (thale cress)
 ORGANISM Arabidopsis thaliana
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
 ; eustosids II; Brassicales; Brassicaceae; Arabidopsis.
 REFERENCE 1 (bases 1 to 334)
 AUTHORS Alonso,J.M., Leisse,T.J., Harajas,P., Chen,H., Cheuk,R., Gadrinab
 ,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.,
 Zimmerman,J. and Ecker,J.K.
 TITLE A Sequence-Indexed Library of Insertion Mutations in the
 Arabidopsis Genome
 JOURNAL Unpublished
 COMMENT Contact: Joseph R. Ecker
 Salk Institute Genomic Analysis Laboratory (SIGNAL)
 The Salk Institute for Biological Studies
 10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
 Tel: 858 453 4100 x1752
 Fax: 858 558 6379
 Email: ecker@salk.edu
 This is single pass sequence recovered from the left border of
 T-DNA.
 Class: T-DNA tagged.
 Location/Qualifiers
 1..334

FEATURES
 source
 1..334
 Location/Qualifiers

```

/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="SALK_082734.31.45.x"
/notes="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at: http://signal.salk.edu/tdna_protocols.html"
BASE COUNT 111 a 71 c 60 g 51 t
ORIGIN

Alignment Scores:
Pred. No.: 480 Length: 334
Score: 41.00 Matches: 7
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 77.78% Mismatches: 0
Query Match: 93.18% Indels: 0
DB: 29 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x B2352875 (1-334)
Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||:|||||:|||||:|||||:
Db 188 TTGTTAGGTGCATTGTGACGAGTATG 162

RESULT 4
CNS000Q2/c
LOCUS
DEFINITION
Arabidopsis thaliana genome survey sequence T7 end of PAC FLOH2 of
IGF library from strain Columbia of Arabidopsis thaliana, genomic
survey sequence.
ACCESSION
AL085272
VERSION
AL085272.1 GI:5286412
KEYWORDS
GSS.
SOURCE
Arabidopsis thaliana (thale cress)
ORGANISM
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.
1 (bases 1 to 400)
Sa.ancubal,M., Chaisne,N., Artiguenave,F., Brottier,P., Wincker,P.,
Samson,D., Saurin,W., Weissbach,J. and Quetier,F.
Unpublished
2 (bases 1 to 400)
Genoscope.
Direct Submission
Submitted (25-JUN-1999) Genoscope - Centre National de Sequence :
BP 191, 91006 Evry cedex - FRANCE (E-mail : seq@genoscope.cns.fr)
Web : www.genoscope.cns.fr
Location/Qualifiers
1..400
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia"
/db_xref="taxon:3702"
/clone="F10H2"
/notes="end : T7"
BASE COUNT 148 a 84 c 61 g 107 t
ORIGIN

Alignment Scores:
Pred. No.: 627 Length: 400
Score: 41.00 Matches: 7
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 77.78% Mismatches: 0
Query Match: 93.18% Indels: 0
DB: 29 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x B2352875 (1-334)
Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||:|||||:|||||:|||||:
Db 188 TTGTTAGGTGCATTGTGACGAGTATG 162

US-09-965-594-1_COPY_14_22 (1-9) x CNS00Q2I (1-400)
Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||:|||||:|||||:|||||:
Db 350 TTGTTAGGTGCATTGTGACGAGTATG 324

RESULT 5
BH758372
LOCUS
DEFINITION
SALK_018581.33.10.x Arabidopsis thaliana TDNA insertion lines
Arabidopsis thaliana genomic clone SALK_018581.33.10.x, genomic
survey sequence.
ACCESSION
BH758372
VERSION
BH758372.1 GI:19043622
KEYWORDS
GSS.
SOURCE
Arabidopsis thaliana (thale cress)
ORGANISM
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eurosids II; Brassicales; Brassicaceae; Arabidopsi.
1 (bases 1 to 424)
Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab
,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.
, Zimmerman,J. and Ecker,J.R.
A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
Unpublished
Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu
This is single pass sequence recovered from the left border of
TDNA.
Class: TDNA tagged.
Location/Qualifiers
1..424
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="SALK_018581.33.10.x"
/notes="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at: http://signal.salk.edu/tdna_protocols.html"
BASE COUNT 62 a 151 c 98 g 113 t
ORIGIN

Alignment Scores:
Pred. No.: 683 Length: 424
Score: 41.00 Matches: 7
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 77.78% Mismatches: 0
Query Match: 93.18% Indels: 0
DB: 28 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x BH758372 (1-424)
Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||:|||||:|||||:|||||:
Db 74 TTGTTAGGTGCATTGTGACGAGTATG 100

RESULT 6
BF651538/c
LOCUS
DEFINITION
274341 MARC 3B0V Bos taurus cDNA 5', rRNA sequence.
ACCESSION
BF651538

```



```

JOURNAL      Unpublished
COMMENT      Other_GSSs: F26N17R
             Contact: Steve Rounsley
             Department of Eukaryotic Genomics
             The Institute for Genomic Research
             9712 Medical Center Dr., Rockville, MD 20850, USA
             Tel: 301 838 0200
             Fax: 301 838 0208
             Email: rounsley@tigr.org
             Seq primer: M13-21
             Class: SAC ends
             High quality sequence stop: 498.
FEATURES     Location/Qualifiers
             ..498
             /organism="Arabidopsis thaliana"
             /mol_type="genomic DNA"
             /strain="Columbia"
             /db_xref="taxon:3702"
             /clone="F26H1"
             /sex="hermaphrodite"
             /note="Vector: pClOACII; Site_1: EcoRI; Site_2: EcoRI;
             Produced by Thomas Altmann"
BASE COUNT   184 a 105 c 80 g 129 t
ORIGIN

Alignment Scores:
Pred. No.:      866      Length:      498
Score:          41.00    Matches:      7
Percent Similarity: 100.00%  Conservative: 2
Best Local Similarity: 77.78%  Mismatches: 0
Query Match:     93.18%  Indels: 0
DB:              28      Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x B96162 (1-498)

Qy 1 LeuLeuGlyCysIlelleThrSerLeu 9
    |||||.....|
Db 370 TTGTTAGGTCGATTGTGACGAGTATG 344

RESULT 9
B26036/c 866 bp DNA linear GSS 13-OCT-1997
LOCUS     B26036
DEFINITION B26036 Arabidopsis thaliana genomic clone F262, genomic survey
sequence.
ACCESSION B26036.1 GI:2512002
VERSION   B26036.1 GI:2512002
KEYWORDS  GSS.
SOURCE    Arabidopsis thaliana (thale cress)
ORGANISM  Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE 1 (bases 1 to 505)
AUTHORS   Rounsley,S.D., Kelley,C.M., Field,C.E., Graven,M.H., Adams,M.D. and
Venter,J.C.
TITLE     Use of a BAC End Sequence Database to Identify Minimal Overlaps for
Arabidopsis Genomic Sequencing
JOURNAL   Unpublished
COMMENT   Contact: Steve Rounsley
             Department of Eukaryotic Genomics
             The Institute for Genomic Research
             9712 Medical Center Dr., Rockville, MT 20850, USA
             Tel: 301 838 0200
             Fax: 301 838 0208
             Email: rounsley@tigr.org
             Seq primer: M13-21
             Class: BAC ends
             High quality sequence stop: 505.
FEATURES     Location/Qualifiers
             1..505
             /organism="Arabidopsis thaliana"

/mol_type="genomic DNA"
/strain="Columbia"
/db_xref="taxon:3702"
/clone="F26H1"
/sex="hermaphrodite"
/clone_lib="JAMJ"
/note="Vector: pClOACII; Site_1: HindIII; Site_2: HindIII
; Produced by Rod Wing"
BASE COUNT   173 a 107 c 93 g 131 t
ORIGIN

Alignment Scores:
Pred. No.:      884      Length:      505
Score:          41.00    Matches:      7
Percent Similarity: 100.00%  Conservative: 2
Best Local Similarity: 77.78%  Mismatches: 0
Query Match:     93.18%  Indels: 0
DB:              28      Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x B26036 (1-505)

Qy 1 LeuLeuGlyCysIlelleThrSerLeu 9
    |||||.....|
Db 196 TTGTTAGGTCGATTGTGACGAGTATG 170

RESULT 10
B2289143/c 651 bp DNA linear GSS 24-OCT-2002
LOCUS     B2289143
DEFINITION SALK_022540.51.60.x Arabidopsis thaliana TDNA insertion lines
Arabidopsis thaliana genomic clone SALK_022540.51.60.x, genomic
survey sequence.
ACCESSION B2289143
VERSION   B2289143.1 GI:24330427
KEYWORDS  GSS.
SOURCE    Arabidopsis thaliana (thale cress)
ORGANISM  Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE 1 (bases 1 to 551)
AUTHORS   Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab
,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Predslis,L., Shinn,P.
,Zimmerman,J. and Ecker,J.R.
TITLE     A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
JOURNAL   Unpublished
COMMENT   Contact: Joseph R. Ecker
             Saik Institute Genomic Analysis Laboratory (SIGAL)
             The Saik Institute for Biological Studies
             10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
             Tel: 858 453 4100 x1752
             Fax: 858 558 6379
             Email: ecker@saik.edu
             This is single pass sequence recovered from the left border of
             TDNA.
             Class: TDNA tagged.
             Location/Qualifiers
             1..651
             /organism="Arabidopsis thaliana"
             /mol_type="genomic DNA"
             /strain="Columbia 0"
             /db_xref="taxon:3702"
             /clone="SALK_022540.51.60.x"
             /clone_lib="Arabidopsis thaliana TDNA insertion lines"
             /note="PCR was performed on Arabidopsis thaliana lines
             each of which contains one or more TDNA insertion
             elements. The resultant fragment for each line was
             directly sequenced to determine the genomic sequence at
             the site of insertion. Details of the protocols used can
             be found at http://signal.salk.edu/tdna_protocols.html"
BASE COUNT   207 a 131 c 113 g 197 t
ORIGIN

```

```

Alignment Scores:
Pred. No.: 1.39e+03 Length: 651
Score: 41.00 Matches: 7
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 77.78% Mismatches: 0
Query Match: 93.18% Indels: 0
DB: 29 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x BH289143 (1-651)
QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||
Db 170 TTGTAGGTGCATTCGACGAGTATG 144

RESULT 11
BH241556 683 bp DNA linear GSS 13-NOV-2001
LOCUS AU1LE777F AUIL Arabidopsis thaliana genomic clone AU1LE77, genomic
survey sequence.
ACCESSION BH241556
VERSION BH241556 GI:16916061
KEYWORDS Arabidopsis thaliana (thale cress)
SOURCE Arabidopsis thaliana
ORGANISM Arabidopsis thaliana
Eukaryota: Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE 1 (bases 1 to 683)
AUTHORS Town,C.D., Whitelaw,C.A., Pai,G., Van Aken,S.E., Utterback,T.V.,
Feldblyum,T.V. and Fraser,C.M.
TITLE Survey sequencing of Arabidopsis thaliana BAC F26J2;
JOURNAL Unpublished
COMMENT Other_GSSs: AU1LE777R
Contact: Chris Town
TIGR
9712 Medical Center Drive, Rockville, MD 20850, USA.
Tel: 301-838-3521
Fax: 301-838-0208
Email: cdtown@tigr.org
From Wash. U contig 720.
Seq primer: TF
Class: sheared ends.
Location/Qualifiers
source
1..683
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia"
/db_xref="taxon:3702"
/clone="AU1577"
/clone_lib="JGI"
/note="Vector: pUC19; Site_1: EcoRI; Site_2: EcoRI;
Produced by Thomas Altmann"
BASE COUNT 196 a 129 c 138 g 220 t
ORIGIN
1..683
1.39e+03 Length: 683
Pred. No.: 41.00 Matches: 7
Score: 100.00% Conservative: 2
Percent Similarity: 77.78% Mismatches: 0
Best Local Similarity: 93.18% Indels: 0
Query Match: 28 Gaps: 0
DB: 0

US-09-965-594-1_COPY_14_22 (1-9) x BH241556 (1-683)
QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||
Db 655 TTGTAGGTGCATTCGACGAGTATG 681

RESULT 12
BH241556 693 bp DNA linear GSS 31-MAR-1998
LOCUS AU1LE777F AUIL Arabidopsis thaliana genomic clone F21D4, genomic
survey sequence.
ACCESSION BH241556
VERSION BH241556 GI:16915962
KEYWORDS Arabidopsis thaliana (thale cress)
SOURCE Arabidopsis thaliana
ORGANISM Arabidopsis thaliana
Eukaryota: Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE 1 (bases 1 to 693)
AUTHORS Town,C.D., Whitelaw,C.A., Pai,G., Van Aken,S.E., Utterback,T.V.,
Feldblyum,T.V. and Fraser,C.M.
TITLE Survey sequencing of Arabidopsis thaliana BAC F26J2;
JOURNAL Unpublished
COMMENT Other_GSSs: AU1LE777R
Contact: Steve Rounsley
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: rounsley@tigr.org
Seq primer: M13-21
Class: BAC ends
High quality sequence stop: 693.
Location/Qualifiers
source
1..693
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia"
/db_xref="taxon:3702"
/clone="F21D4"
/sex="hermaphrodite"
/clone_lib="JGI"
/note="Vector: pUC19; Site_1: EcoRI; Site_2: EcoRI;
Produced by Thomas Altmann"
BASE COUNT 210 a 117 c 144 g 221 t
ORIGIN
1..693
1.41e+03 Length: 693
Pred. No.: 41.00 Matches: 7
Score: 100.00% Conservative: 2
Percent Similarity: 77.78% Mismatches: 0
Best Local Similarity: 93.18% Indels: 0
Query Match: 28 Gaps: 0
DB: 0

US-09-965-594-1_COPY_14_22 (1-9) x BH241556 (1-693)
QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||
Db 480 TTGTAGGTGCATTCGACGAGTATG 506

RESULT 13
BH241494 719 bp DNA linear GSS 13-NOV-2001
LOCUS AU1LF221R AUIL Arabidopsis thaliana genomic clone AU1LF22, genomic
survey sequence.
ACCESSION BH241494
VERSION BH241494 GI:16915962
KEYWORDS Arabidopsis thaliana (thale cress)
SOURCE Arabidopsis thaliana
ORGANISM Arabidopsis thaliana
Eukaryota: Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE 1 (bases 1 to 719)
AUTHORS Town,C.D., Whitelaw,C.A., Pai,G., Van Aken,S.E., Utterback,T.V.,
Feldblyum,T.V. and Fraser,C.M.
TITLE Survey sequencing of Arabidopsis thaliana BAC F26J2;
JOURNAL Unpublished
COMMENT Other_GSSs: AU1LF221R

```

Contact: Chris Town

TIGR
9712 Medical Center Drive, Rockville, MD 20850, USA.

Tel: 301-838-3523

Fax: 301-838-0208

Email: cdtown@tigr.org

From Wash. U contig 720.

Seq primer: TF

Class: Sheared ends.

Location/Qualifiers

1..719

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/strain="Columbia"

/db_xref="taxon:3702"

/clone_lib="AULF82"

/clone_lib="AULF"

/note="Vector: pPOS2; Site 1: BstXI; 2-3 kb sheared BAC DNA inserted into pPOS2 using BstXI linkers"

197 a 149 c 140 g 234 t

BASE COUNT

ORIGIN

Alignment Scores:

Pred. No.: 1,49e-03 Length: 719

Score: 41.00 Matches: 7

Percent Similarity: 100.00% Conservatives: 2

Best Local Similarity: 77.78% Mismatches: 0

Query Match: 93.18% Indels: 0

DB: 28 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x BH241494 (1-719)

Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9

Db 130 TTGTTAGTTCATTGTGACGAGTATG 156

RESULT 14

LOCUS

DEFINITION LERAD347F LERA Arabidopsis thaliana genomic clone LERAD347, genomic survey sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Arabidopsis thaliana (thale cress)

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids

; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE

AUTHORS

1 (bases 1 to 724)

Rue, C.R., Lin, X., Pai, G., Barnstead, M., Bowman, C., Utterback, T.,

Feldblyum, J., Liang, F., Creasy, T., and Fraser, C.M.

Genomic survey sequencing of Landsberg erecta ecotype 01

Arabidopsis thaliana and identification of sequence-based

polymorphisms

unpublished

Contact: Xiaoying Lin

The Institute for Genomic Research

9712 Medical Center Dr., Rockville, MD 20850, USA

Tel: 301 838 0200

Fax: 301 838 0208

Email: at@tigr.org

Seq primer: TF

Class: shotgun.

Location/Qualifiers

1..724

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/strain="Landsberg erecta"

/db_xref="taxon:3702"

/clone="LERAD34"

/clone_lib="LERA"

FEATURES

source

/note="Organ: Leaf; Vector: pPOS1; total genomic DNA was sheared to 0.9-1 Kbp before ligation."

187 a 151 c 143 g 243 t

BASE COUNT

ORIGIN

Alignment Scores:

Pred. No.: 1,5e+03 Length: 724

Score: 41.00 Matches: 7

Percent Similarity: 100.00% Conservatives: 2

Best Local Similarity: 77.78% Mismatches: 0

Query Match: 93.18% Indels: 0

DB: 28 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AQ955329 (1-724)

Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9

Db 70 TTGTTAGTTCATTGTGACGAGTATG 96

RESULT 15

LOCUS

DEFINITION BH241165 Arabidopsis thaliana genomic clone AULF85, genomic survey sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Arabidopsis thaliana (thale cress)

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids

; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE

AUTHORS

1 (bases 1 to 758)

Town, C.D., Whitelaw, C.A., Pai, G., Van Aken, S.E., Utterback, T.V.,

Feldblyum, T.V. and Fraser, C.M.

Survey sequencing of Arabidopsis thaliana BAC F25J21

unpublished

Other GSSs: AULF85TR

Contact: Chris Town

TIGR

9712 Medical Center Drive, Rockville, MD 20850, USA.

Tel: 301-838-3523

Fax: 301-838-0208

Email: cdtown@tigr.org

From Wash. U contig 720.

Seq primer: TF

Class: sheared ends.

Location/Qualifiers

1..758

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/strain="Columbia"

/db_xref="taxon:3702"

/clone="AULF85"

/clone_lib="AULF"

/note="Vector: pPOS2; Site 1: BstXI; 2-3 kb sheared BAC DNA inserted into pPOS2 using BstXI linkers"

262 a 140 c 157 g 259 t

BASE COUNT

ORIGIN

Alignment Scores:

Pred. No.: 1,61e+03 Length: 758

Score: 41.00 Matches: 7

Percent Similarity: 100.00% Conservatives: 2

Best Local Similarity: 77.78% Mismatches: 0

Query Match: 93.18% Indels: 0

DB: 28 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x BH241165 (1-758)

Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9

Db 318 TTGTTAGTTCATTGTGACGAGTATG 344

Search completed: September 29, 2003, 20:22:55
Job time : 1632 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 Computer Ltd.

OM protein - protein search, using sw model

Run on: September 23, 2003, 19:00:07 : Search time 30 Seconds
(without alignments)
77,416 Million cell updates/sec

Title: US-09-965-594-1_COPY_14_22

Perfect score: 44

Sequence: 1 LLAGCIITSL 9

Scoring table: BLOSUM62

Gapop 13.0, Gapext 0.5

Searched: 830525 seqs, 25952694 residues

Total number of hits satisfying chosen parameters: 830296

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 99%

Listing first 45 summaries

Database :

- 1: SP_ARCHAEA*
- 2: SP_BACTERIA*
- 3: SP_FUNGI*
- 4: SP_HUMAN*
- 5: SP_INVERTEBRATE*
- 6: SP_MAMMAL*
- 7: SP_MMC*
- 8: SP_ORGANELLAE*
- 9: SP_PHAGE*
- 10: SP_PLANT*
- 11: SP_RODENT*
- 12: SP_VIRUS*
- 13: SP_VERTEBRATE*
- 14: SP_UNCLASSIFIED*
- 15: SP_VIRUS*
- 16: SP_BACTERIAP*
- 17: SP_ARCHAEA*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	43	97.7	180	12	Q8QW28
2	43	97.7	180	12	Q8QW29
3	43	97.7	180	12	Q8QW27
4	43	97.7	180	12	Q8QW23
5	43	97.7	403	12	Q8QW94
6	43	97.7	3010	12	Q9QIY5
7	43	97.7	3010	12	Q9QIY6
8	43	97.7	3010	12	Q9QIY2
9	43	97.7	3010	12	Q8V638
10	43	97.7	3010	12	Q8V638
11	43	97.7	3010	12	Q9QIY6
12	43	97.7	3011	12	Q9I304
13	42	95.5	181	12	Q9I3P5
14	42	95.5	3010	12	Q9J3B9
15	42	95.5	3010	12	Q9J3C5
16	41	93.2	180	12	Q8QW19

17	41	93.2	361	12	Q70817	Q70817 hepatitis C
18	41	93.2	2864	12	Q92973	Q92973 hepatitis C
19	41	93.2	2864	12	Q9W1K8	Q9W1K8 hepatitis C
20	41	93.2	2864	12	Q92976	Q92976 hepatitis C
21	41	93.2	3010	12	Q02829	Q02829 hepatitis C
22	41	93.2	3010	12	Q92959	Q92959 hepatitis C
23	41	93.2	3010	12	Q02828	Q02828 hepatitis C
24	41	93.2	3010	12	Q9J3G0	Q9J3G0 hepatitis C
25	40	90.9	180	12	Q8QVX2	Q8QVX2 hepatitis C
26	40	90.9	180	12	Q8QW22	Q8QW22 hepatitis C
27	40	90.9	180	12	Q8QW09	Q8QW09 hepatitis C
28	40	90.9	180	12	Q8QW26	Q8QW26 hepatitis C
29	40	90.9	180	12	Q8QVW8	Q8QVW8 hepatitis C
30	40	90.9	180	12	Q8QVX0	Q8QVX0 hepatitis C
31	40	90.9	180	12	Q8QVW2	Q8QVW2 hepatitis C
32	40	90.9	180	12	Q8QVZ2	Q8QVZ2 hepatitis C
33	40	90.9	180	12	Q8QVZ6	Q8QVZ6 hepatitis C
34	40	90.9	181	12	Q9I3Q7	Q9I3Q7 hepatitis C
35	40	90.9	181	12	Q9I3Q6	Q9I3Q6 hepatitis C
36	40	90.9	2864	12	Q9W1K9	Q9W1K9 hepatitis C
37	40	90.9	2864	12	Q9W1L0	Q9W1L0 hepatitis C
38	40	90.9	2864	12	Q92975	Q92975 hepatitis C
39	40	90.9	3010	12	Q90192	Q90192 hepatitis C
40	40	90.9	3010	12	Q9C1Y7	Q9C1Y7 hepatitis C
41	40	90.9	3010	12	Q92970	Q92970 hepatitis C
42	40	90.9	3010	12	Q9J3G9	Q9J3G9 hepatitis C
43	40	90.9	3010	12	Q9J3G4	Q9J3G4 hepatitis C
44	40	90.9	3010	12	Q9QIY8	Q9QIY8 hepatitis C
45	40	90.9	3010	12	Q9QIY8	Q9QIY8 hepatitis C

ALIGNMENTS

RESULT 1

Q8QW08 ID Q8QW08 PRELIMINARY: PRI: 180 AA.

AC Q8QW08; 01-JUN-2002 (TrEMBLrel. 21, Created)
 DJ Q8QW08; 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
 DI Q8QW08; 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
 DE Polypeptide (Fragment).

CS Hepatitis C virus type 1b.
 OC Viruses; ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
 OC Hepacivirus.

CX NCBI_TaxID=31647;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=108;

RA Qdata S.;

RT *CORRELATION BETWEEN SECONDARY STRUCTURE OF AN AMINO-TERMINAL PORTION OF THE NONSTRUCTURAL PROTEIN 3 OF HEPATITIS C VIRUS AND DEVELOPMENT OF HEPATOCELLULAR CARCINOMA.*

RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.

DR EMBL; AB072067; BA588249.1; -

DR InterPro; IPR004109; HCV_NS3.

DR Pfam; PF02907; HCV_NS3; 1.

FT NON_TER 1

FT NON_TER 180

SQ SEQUENCE 180 AA: 18982 MW: 77E64C2673D89DEC CRC64;

Query Match 97.7%; Score 43; DB 12; Length 180;

Best Local Similarity 88.9%; Pred. No. 1.1;

Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 1 LLAGCIITSL 9

Db 13 LLAGCIITSL 21

RESULT 2

Q8QW28 ID Q8QW28 PRELIMINARY: PRI: 180 AA.

```

AC Q8QW28;
DI 01-JUN-2002 (TrEMBLrel. 21, Created)
DI 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
DI 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Polyprotein (Fragment).
OS Hepatitis C virus type 1b.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus
OX NCBI_TaxID=31647;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=H05-4;
RA Oyata S.;
RT "CORRELATION BETWEEN SECONDARY STRUCTURE OF AN AMINO-TERMINAL PORTION
RT OF THE NONSTRUCTURAL PROTEIN 3 OF HEPATITIS C VIRUS AND DEVELOPMENT OF
RT HEPATOCELLULAR CARCINOMA.";
RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB072047; BAB88229.1; -.
DR InterPro; IPR004109; HCV_NS3.
DR Pfam; PF02907; HCV_NS3; 1.
FT NON_TER 1 180
FT NON_TER 180 180
SQ SEQUENCE 180 AA: 18891 MW: 85AFSPSELCF2BCA30 CRC64;

Query Match 97.7%; Score 43; DB 12; Length 180;
Best Local Similarity 88.9%; Pred. No. 1.1;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LLGCIITSL 9
DB 13 LLGCIIVTSL 21

RESULT 3
Q8QW27
ID Q8QW27 PRELIMINARY; PRT; 180 AA.
AC Q8QW27;
DI 01-JUN-2002 (TrEMBLrel. 21, Created)
DI 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
DI 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Polyprotein (Fragment).
OS Hepatitis C virus type 1b.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus
OX NCBI_TaxID=31647;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=H05-5;
RA Oyata S.;
RT "CORRELATION BETWEEN SECONDARY STRUCTURE OF AN AMINO-TERMINAL PORTION
RT OF THE NONSTRUCTURAL PROTEIN 3 OF HEPATITIS C VIRUS AND DEVELOPMENT OF
RT HEPATOCELLULAR CARCINOMA.";
RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB072048; BAB88230.1; -.
DR InterPro; IPR004109; HCV_NS3.
DR Pfam; PF02907; HCV_NS3; 1.
FT NON_TER 1 180
FT NON_TER 180 180
SQ SEQUENCE 180 AA: 18913 MW: 895FB21253DA6CD6 CRC64;

Query Match 97.7%; Score 43; DB 12; Length 180;
Best Local Similarity 88.9%; Pred. No. 1.1;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LLGCIITSL 9
DB 13 LLGCIIVTSL 21

RESULT 4
Q8QVX3
ID Q8QVX3 PRELIMINARY; PRT; 180 AA.
AC Q8QVX3;

```

```

LT 01-JUN-2002 (TrEMBLrel. 21, Created)
DI 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
DI 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Polyprotein (Fragment).
OS Hepatitis C virus type 1b.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus
OX NCBI_TaxID=31647;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=N-82;
RA Oyata S.;
RT "CORRELATION BETWEEN SECONDARY STRUCTURE OF AN AMINO-TERMINAL PORTION
RT OF THE NONSTRUCTURAL PROTEIN 3 OF HEPATITIS C VIRUS AND DEVELOPMENT OF
RT HEPATOCELLULAR CARCINOMA.";
RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB072102; BAB88284.1; -.
DR InterPro; IPR004109; HCV_NS3.
DR Pfam; PF02907; HCV_NS3; 1.
FT NON_TER 1 180
FT NON_TER 180 180
SQ SEQUENCE 180 AA: 18808 MW: CE520CF0BCA4E1E2 CRC64;

Query Match 97.7%; Score 43; DB 12; Length 180;
Best Local Similarity 88.9%; Pred. No. 1.1;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LLGCIITSL 9
DB 13 LLGCIIVTSL 21

RESULT 5
Q8QF94
ID Q8QF94 PRELIMINARY; PRT; 403 AA.
AC Q8QF94;
DI 01-JUN-2002 (TrEMBLrel. 21, Created)
DI 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
DI 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Genome polyprotein (Fragment).
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=667;
RX MEDLINE=21904745; PubMed=1197242;
RA Kallings O., Norder H., Mukomolov S., Magnus L.O.;
RT "A natural intergenotypic recombinant of hepatitis C virus identified
RT in St. Petersburg.";
PL J. Virol. 76:4034-4043(2002).
DR EMBL; AY070171; AAL58587.1; -.
DR InterPro; IPR02531; HCV_NS1.
DR InterPro; IPR02518; HCV_NS2.
DR InterPro; IPR004109; HCV_NS3.
DR Pfam; PF01560; HCV_NS1; 1.
DR Pfam; PF01538; HCV_NS2; 1.
DR Pfam; PF02907; HCV_NS3; 1.
KW Coat protein; Envelope protein; Glycoprotein; Nonstructural protein;
FT NON_TER 1 403
FT NON_TER 403 403
SQ SEQUENCE 403 AA: 43868 MW: EASC9FACB9348C8A CRC64;

Query Match 97.7%; Score 43; DB 12; Length 403;
Best Local Similarity 88.9%; Pred. No. 2.2;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LLGCIITSL 9
DB 323 LLGCIIVTSL 331

```

RESULT 6

Q90IY5 PRELIMINARY: PRT: 3010 AA.
 AC Q90IY5: (1)
 DT 01-MAY-2000 (TRENBLREL. 23, Created)
 DI 01-MAY-2000 (TRENBLREL. 23, Last sequence update)
 DE 01-MAR-2003 (TRENBLREL. 23, Last annotation update)
 DE Genome polyprotein.
 OS Hepatitis C virus.
 CC Viruses: ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
 CC Hepacivirus.
 CC NCBI_TaxID=11103;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MD3-2;
 RX MEDLINE=20013325; PubMed=12544098;
 RA Nagayama K., Kurosaki M., Enomoto N., Maekawa S.Y., Miyasaka Y.,
 RA Tazawa J.I., Izumi N., Marumo F., Sato C.;
 RT "Time-related changes in full-length hepatitis C virus and hepatitis
 RT activity.";
 RL Virology 263:244-253(1999).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MD3-2;
 RA Nagayama K., Kurosaki M., Enomoto N., Maekawa S., Miyasaka Y.,
 RA Sakamoto N., Fukuma T., Iizawa J., Izumi N., Marumo F., Sato C.;
 RL Submitted (JUL-1999) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
 CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
 CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
 CC PROTEIN C AND MRNA (BY SIMILARITY).
 CC EMBL: AF155050; AAD56185.1; .
 DR HSP: P26563; IJXP.
 DR InterPro: IPR001410; DEAD.
 DR InterPro: IPR002522; HCV_capsid.
 DR InterPro: IPR002521; HCV_core.
 DR InterPro: IPR002519; HCV_env.
 DR InterPro: IPR002531; HCV_NS1.
 DR InterPro: IPR002518; HCV_NS2.
 DR InterPro: IPR004109; HCV_NS3.
 DR InterPro: IPR000745; HCV_NS4a.
 DR InterPro: IPR001490; HCV_NS4b.
 DR InterPro: IPR002868; HCV_NS5a.
 DR InterPro: IPR002166; HCV_RDRP.
 DR InterPro: IPR007095; RNA_pol_PS.
 DR InterPro: IPR007034; RNA_pol_PSVir.
 DR Pfam: PF01543; HCV_capsid; 1.
 DR Pfam: PF01542; HCV_core; 1.
 DR Pfam: PF01539; HCV_env; 1.
 DR Pfam: PF01560; HCV_NS1; 1.
 DR Pfam: PF01538; HCV_NS2; 1.
 DR Pfam: PF02907; HCV_NS3; 1.
 DR Pfam: PF01206; HCV_NS4a; 1.
 DR Pfam: PF01901; HCV_NS4b; 1.
 DR Pfam: PF01506; HCV_NS5a; 1.
 DR Pfam: PF00998; Viral_RDRP; 1.
 DR ProDom: PD186062; HCV_NS1; 1.
 DR SMART: SM00487; DEXDC; 1.
 DR PROSITE: PS05057; RDRP_POSITIVE; 1.
 DR PROSITE: PS05052; RDRP_VIRAL; 1.
 KW Coat protein: Envelope protein: Glycoprotein: Nonstructural protein;
 KW Polyprotein: RNA-directed RNA polymerase: Transferase: Transmembrane.
 SQ SEQUENCE 3010 AA: 327430 MW: 15190E3463DRAC15 CRC64;

Query Match 97.7% Score 43; DB 12; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 12;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LIGCIITSL 9
 :|||||:
 Db 1039 LIGCIVTSL 1047

RESULT 7

Q90IY6 PRELIMINARY: PRT: 3010 AA.
 AC Q90IY6: (1)
 DT 01-MAY-2000 (TRENBLREL. 13, Created)
 DI 01-MAY-2000 (TRENBLREL. 13, Last sequence update)
 DE 01-MAR-2003 (TRENBLREL. 23, Last annotation update)
 DE Genome polyprotein.
 OS Hepatitis C virus.
 CC Viruses: ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
 CC Hepacivirus.
 CC NCBI_TaxID=11103;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MD3-1;
 RX MEDLINE=20013325; PubMed=10544098;
 RA Nagayama K., Kurosaki M., Enomoto N., Maekawa S.Y., Miyasaka Y.,
 RA Tazawa J.I., Izumi N., Marumo F., Sato C.;
 RT "Time-related changes in full-length hepatitis C virus and hepatitis
 RT activity.";
 RL Virology 263:244-253(1999).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MD3-1;
 RA Nagayama K., Kurosaki M., Enomoto N., Maekawa S., Miyasaka Y.,
 RA Sakamoto N., Fukuma T., Iizawa J., Izumi N., Marumo F., Sato C.;
 RL Submitted (JUL-1999) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
 CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
 CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
 CC PROTEIN C AND MRNA (BY SIMILARITY).
 CC EMBL: AF155049; AAD56184.1; .
 DR HSP: P26663; IJXP.
 DR InterPro: IPR001410; DEAD.
 DR InterPro: IPR002522; HCV_capsid.
 DR InterPro: IPR002521; HCV_core.
 DR InterPro: IPR002519; HCV_env.
 DR InterPro: IPR002531; HCV_NS1.
 DR InterPro: IPR002518; HCV_NS2.
 DR InterPro: IPR004109; HCV_NS3.
 DR InterPro: IPR000745; HCV_NS4a.
 DR InterPro: IPR001490; HCV_NS4b.
 DR InterPro: IPR002868; HCV_NS5a.
 DR InterPro: IPR002166; HCV_RDRP.
 DR InterPro: IPR007095; RNA_pol_DS_PS.
 DR InterPro: IPR007034; RNA_pol_PSVir.
 DR Pfam: PF01543; HCV_capsid; 1.
 DR Pfam: PF01542; HCV_core; 1.
 DR Pfam: PF01539; HCV_env; 1.
 DR Pfam: PF01560; HCV_NS1; 1.
 DR Pfam: PF01538; HCV_NS2; 1.
 DR Pfam: PF02907; HCV_NS3; 1.
 DR Pfam: PF01066; HCV_NS4a; 1.
 DR Pfam: PF01001; HCV_NS4b; 1.
 DR Pfam: PF01506; HCV_NS5a; 1.
 DR Pfam: PF00998; Viral_RDRP; 1.
 DR ProDom: PD186062; HCV_NS1; 1.
 DR SMART: SM00487; DEXDC; 1.
 DR PROSITE: PS05057; RDRP_POSITIVE; 1.
 DR PROSITE: PS05052; RDRP_VIRAL; 1.
 KW Coat protein: Envelope protein: Glycoprotein: Nonstructural protein;
 KW Polyprotein: RNA-directed RNA polymerase: Transferase: Transmembrane.
 SQ SEQUENCE 3010 AA: 327368 MW: 998C7F293EAAEC8D CRC64;

Query Match 97.7% Score 43; DB 12; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 12;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LIGCIITSL 9
 :|||||:
 Db 1039 LIGCIVTSL 1047

01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
Genome polyprotein.
Hepatitis C virus.
Viruses: ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepadnaviridae;
NCBI_TaxID=11103;
[1];
SEQUENCE FROM N.A.
STRAIN=J1;
MEDLINE=92295714; PubMed=1318627;
Tanaka T., Kato N., Nakagawa M., Gotsuyama Y., Cho M.J., Nakazawa I.,
Hijikata M., Ishimura Y., Shimotohno K.;
*Molecular cloning of hepatitis C virus genome from a single Japanese
carrier: sequence variation within the same individual and among
infected individuals*;
Virus Res. 23:39-54(1992).
-1- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
PROTEIN C AND RNA (BY SIMILARITY).
EMBL: D11355; BAAL8894.1;
HSSP: P26563; IJXP.
InterPro: IPR001410; DEAD.
InterPro: IPR002522; HCV_capsid.
InterPro: IPR002522; HCV_core.
InterPro: IPR002519; HCV_env.
InterPro: IPR002533; HCV_NS1.
InterPro: IPR002518; HCV_NS2.
InterPro: IPR004109; HCV_NS3.
InterPro: IPR000745; HCV_NS4a.
InterPro: IPR001490; HCV_NS4b.
InterPro: IPR002868; HCV_NS5a.
InterPro: IPR002156; HCV_NS5b.
InterPro: IPR007095; RNA_pol_DS_PS.
InterPro: IPR007094; RNA_pol_PSVir.
Pfam: PF01543; HCV_core; 1.
Pfam: PF01542; HCV_core; 1.
Pfam: PF01539; HCV_env; 1.
Pfam: PF01560; HCV_NS1; 1.
Pfam: PF01538; HCV_NS2; 1.
Pfam: PF01538; HCV_NS2; 1.
Pfam: PF02907; HCV_NS3; 1.
Pfam: PF01006; HCV_NS4a; 1.
Pfam: PF01001; HCV_NS4b; 1.
Pfam: PF01506; HCV_NS5a; 1.
Pfam: PF00998; Viral_RdRP; 1.
ProDom: PD186062; HCV_NS1; 1.
SMART: SM00487; DEXDC; 1.
PROSITE: PS00150; CYTOCHROME_C; 1.
PROSITE: PS05507; RDRP_POSITIVE; 1.
PROSITE: PS05521; RDRP_VIRAL; 1.
Coat protein; Envelope protein; Glycoprotein; Nonstructural protein;
Polyprotein; RNA-directed RNA polymerase; Transferrase; Transmembrane;
FT CHAIN 1 191
FT CHAIN 192 383
FT CHAIN 384 729
FT CHAIN 730 1006
FT CHAIN 1007 1615
FT CHAIN 1616 1862
FT CHAIN 1863 2013
FT CHAIN 2014 3010
SEQUENCE 3010 AA; 326564 MK; C5F0B21C2F5D09D CRC64;
Query Match 97.7%; Score 43; DR 12; Length 3010;
Best Local Similarity 88.9%; Pred. No. 12;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 LIGCITITSL 9
D6 1039 LLGCVITSL 1047
RESULT 11
Q913D6 PRELIMINARY; PRT: 3010 AA.
ID Q913D6

Q913D6;
01-MAR-2001 (TrEMBLrel. 16, Created)
01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
Genome polyprotein.
Viruses: ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepadnaviridae;
NCBI_TaxID=11103;
[1];
SEQUENCE FROM N.A.
STRAIN=HCV2221;
Takahashi K., Iwata K., Matsumoto M., Matsumoto H., Nakao K.,
Hatahara I., Ohta Y., Kanai K., Maruo H., Baba K., Hijikata M.,
Watanabe S.;
*Hepatitis C virus (HCV) genotype 1b sequences from fifteen patients
with hepatocellular carcinoma: the 'progression score' revisited*;
Submitted (SEP-2000) to the EMBL/GenBank/DBJ databases.
-1- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
PROTEIN C AND RNA (BY SIMILARITY).
EMBL: AB049101; BAB18814.1;
HSSP: P26563; IJXP.
InterPro: IPR003345; CytC_heme_bind.
InterPro: IPR001410; DEAD.
InterPro: IPR002522; HCV_capsid.
InterPro: IPR002531; HCV_core.
InterPro: IPR002519; HCV_env.
InterPro: IPR002531; HCV_NS1.
InterPro: IPR002518; HCV_NS2.
InterPro: IPR004109; HCV_NS3.
InterPro: IPR000745; HCV_NS4a.
InterPro: IPR001490; HCV_NS4b.
InterPro: IPR002868; HCV_NS5a.
InterPro: IPR002166; HCV_NS5b.
InterPro: IPR001650; Helicase_C.
InterPro: IPR007095; RNA_pol_DS_PS.
InterPro: IPR007094; RNA_pol_PSVir.
Pfam: PF01543; HCV_capsid; 1.
Pfam: PF01542; HCV_core; 1.
Pfam: PF01539; HCV_env; 1.
Pfam: PF01560; HCV_NS1; 1.
Pfam: PF01538; HCV_NS2; 1.
Pfam: PF02907; HCV_NS3; 1.
Pfam: PF01006; HCV_NS4a; 1.
Pfam: PF01001; HCV_NS4b; 1.
Pfam: PF01506; HCV_NS5a; 1.
Pfam: PF00271; Helicase_C; 1.
Pfam: PF00998; Viral_RdRP; 1.
ProDom: PD186062; HCV_NS1; 1.
SMART: SM00487; DEXDC; 1.
PROSITE: PS00150; CYTOCHROME_C; 1.
PROSITE: PS05507; RDRP_POSITIVE; 1.
PROSITE: PS05521; RDRP_VIRAL; 1.
ATP-binding; Coat protein; Envelope protein; Glycoprotein; Helicase;
Hydrolase; Nonstructural protein; Polyprotein;
RNA-directed RNA polymerase; Transferrase; Transmembrane;
SEQUENCE 3010 AA; 327108 MK; DE182D810EF78E54 CRC64;
Query Match 97.7%; Score 43; DB 12; Length 3010;
Best Local Similarity 88.9%; Pred. No. 12;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 LIGCITITSL 9
D6 1039 LLGCVITSL 1047
RESULT 12
Q913D4 PRELIMINARY; PRT: 3011 AA.
ID Q913D4
AC Q913D4

DT 01-DEC-2001 (TrEMBLrel. 19, Created;
 DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE Genome polyprotein.
 OS Hepatitis C virus.
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 OC Hepacivirus.
 OX NCBI_TaxID=11103;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Guntaka K.V., Kuppali S.K., Khaja M.N., Kota K.K., Ramana V.K.,
 RA Swaminathan S., Sakata Y., Habeebullah C.M.;
 RT "Nucleotide Sequence of Indian strain of Hepatitis C Virus."
 RL Submitted (AUG-2001) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
 CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
 CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
 CC PROTEIN C AND MRNA (BY SIMILARITY).
 DR EMBL: AY051292; AAK95932.1; -;
 DR MEROPS: S29.001; -;
 DR MEROPS: U39.001; -;
 DR InterPro: IPR000345; CytC_heme_bind.
 DR InterPro: IPR001410; DEAD.
 DR InterPro: IPR002522; HCV_capsid.
 DR InterPro: IPR002521; HCV_core.
 DR InterPro: IPR002519; HCV_env.
 DR InterPro: IPR002531; HCV_NS1.
 DR InterPro: IPR002518; HCV_NS2.
 DR InterPro: IPR004109; HCV_NS3.
 DR InterPro: IPR000745; HCV_NS4a.
 DR InterPro: IPR001490; HCV_NS4b.
 DR InterPro: IPR002868; HCV_NS5a.
 DR InterPro: IPR032166; HCV_RdRP.
 DR InterPro: IPR007095; RNA_pol_DS_PS.
 DR InterPro: IPR037694; RNA_pol_PSVir.
 DR Pfam: PF01543; HCV_capsid; 1.
 DR Pfam: PF01542; HCV_core; 1.
 DR Pfam: PF01560; HCV_NS1; 1.
 DR Pfam: PF02907; HCV_NS3; 1.
 DR Pfam: PF01006; HCV_NS4a; 1.
 DR Pfam: PF01001; HCV_NS4b; 1.
 DR Pfam: PF01506; HCV_NS5a; 1.
 DR Pfam: PF00398; Viral_RdRP; 1.
 DR ProDom: PD186062; HCV_NS1; 1.
 DR SMART: SM00487; DEXDC; 1.
 DR PROSITE: PS00190; CYTOCHROME_C; 1.
 DR PROSITE: PS0507; RDRP_POSITIVE; 1.
 DR PROSITE: PS0521; RDRP_VIRAL; 1.
 KW Coat protein: envelope protein; Glycoprotein: Nonstructural protein.
 KW Polyprotein: RNA-directed RNA polymerase; Transcriptase; Transmembrane.
 SQ SEQUENCE 3011 AA: 327234 MW: 57A21963A422785C CRC64;

 Query Match 97.7%; Score 43; DB 12; Length 3011;
 Best Local Similarity 88.9%; Pred. No. 12;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

 QY 1 LLGCIITSL 9
 Db 1039 LLGCIIVTSL 1047

 RESULT 13
 Q91RP5
 ID Q91RP5 PRELIMINARY: PRT; 181 AA.
 AC Q91RP5
 DT 01-DEC-2001 (TrEMBLrel. 19, Created)
 DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE NS3 protease (fragment).
 OS Hepatitis C virus.
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;

OC Hepacivirus.
 OX NCBI_TaxID=11103;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Holland-Staley C.A., Kovari L.C., Golenberg E., Meyers D.L.;
 RT "Genetic Diversity and Response to IFN of the NS3 Protease Gene from
 RT Clinical Strains of the Hepatitis C Virus."
 RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF369258; AAK54583.1; -;
 DR MEROPS: S29.001; -;
 DR MEROPS: U39.001; -;
 DR InterPro: IPR004109; HCV_NS3.
 DR Pfam: PF02907; HCV_NS3; 1.
 KW Protease.
 FT NON_TER 1 181
 FT NON_TER 181 181
 FT NON_TER 181 181
 SQ SEQUENCE 181 AA: 19056 MW: BAF89690AD2971DB CRC64;

 Query Match 95.5%; Score 42; DB 12; Length 181;
 Best Local Similarity 88.9%; Pred. No. 1.8;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

 QY 1 LLGCIITSL 9
 Db 13 LLGCIITSL 21

 RESULT 14
 Q9J3H9
 ID Q9J3H9 PRELIMINARY: PRT; 3010 AA.
 AC Q9J3H9
 DT 01-OCT-2000 (TrEMBLrel. 15, Created)
 DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE Genome polyprotein.
 OS Hepatitis C virus.
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 OC Hepacivirus.
 OX NCBI_TaxID=11103;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Nagayama K., Kurosaki M., Enomoto N., Miyasaka Y., Marumoto F., Sato C.;
 RT "Characteristics of hepatitis C virus genome associated with disease
 RT progression."
 RL Submitted (NOV-1999) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
 CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
 CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
 CC PROTEIN C AND MRNA (BY SIMILARITY).
 DR EMBL: AF207734; AAF65944.1; -;
 DR HSSP: F26663; IUXP.
 DR InterPro: IPR000345; CytC_heme_bind.
 DR InterPro: IPR001410; DEAD.
 DR InterPro: IPR002522; HCV_capsid.
 DR InterPro: IPR002521; HCV_core.
 DR InterPro: IPR002519; HCV_env.
 DR InterPro: IPR002531; HCV_NS1.
 DR InterPro: IPR002518; HCV_NS2.
 DR InterPro: IPR004109; HCV_NS3.
 DR InterPro: IPR000745; HCV_NS4a.
 DR InterPro: IPR001490; HCV_NS4b.
 DR InterPro: IPR002868; HCV_NS5a.
 DR InterPro: IPR002156; HCV_RdRP.
 DR InterPro: IPR007095; RNA_pol_DS_PS.
 DR InterPro: IPR007094; RNA_pol_PSVir.
 DR Pfam: PF01543; HCV_capsid; 1.
 DR Pfam: PF01542; HCV_core; 1.
 DR Pfam: PF01539; HCV_env; 1.
 DR Pfam: PF01560; HCV_NS1; 1.
 DR Pfam: PF01538; HCV_NS2; 1.
 DR Pfam: PF02907; HCV_NS3; 1.

```

DR Pfam: PF01006; HCV_NS4a; 1.
DR Pfam: PF01001; HCV_NS4b; 1.
DR Pfam: PF01506; HCV_NS5a; 1.
DR Pfam: PF00998; Viral_RdRp; 1.
DR ProDom: PD186062; HCV_NS1; 1.
DR SMART: SMO0487; DEXDC; 1.
DR PROSITE: PS00190; CYTOCHROME_C; 1.
DR PROSITE: PS05057; RDRP_POSITIVE; 1.
DR PROSITE: PS05052; RDRP_VIRAL; 1.
DR ProDom: PD186062; DEXDC; 1.
DR PROSITE: PS00190; CYTOCHROME_C; 1.
DR PROSITE: PS05057; RDRP_POSITIVE; 1.
DR PROSITE: PS05052; RDRP_VIRAL; 1.
KW Coat protein; Envelope protein; Glycoprotein; Nonstructural protein;
KW Polyprotein; RNA-directed RNA polymerase; Transferase; Transmembrane.
SQ SEQUENCE 3010 AA; 326984 MW; AF120606048078 CR664;

Query Match 95.5%; Score 42; DB 12; Length 3010;
Best Local Similarity 86.9%; Pred. No. 19;
Matches 8; Conservativity 1; Mismatches 6; Indels 0; Gaps 0;

QY 1 LIGCITSL 9
DB 1039 MLCITSL 1047

RESULT 15
Q9J3G5 PRELIMINARY: PRG: 3010 AA.
AC Q9J3G5;
DT 01-OCT-2003 (TRENHirel_15, Created)
DI 01-OCT-2003 (TRENHirel_15, Last sequence update)
DI 01-MAR-2003 (TRENHirel_23, Last annotation update)
DE Genome polyprotein.
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID:11103;
RN [1]
RC SEQUENCE FROM N.A.
RT STRAIN-MD27;
RA Nagayama K., Kurosaki M., Enomoto N., Miyasaka Y., Marumo F., Sato C.;
RT *Characteristics of hepatitis C viral genome associated with disease
RT progression*.
RL Submitted (NOV-1999) to the EMBL/GenBank/DBJ databases.
CC -!- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
CC PROTEIN C AND MRNA (BY SIMILARITY).
DR EMBL: AF207768; AAF65958.1;
DR HSP; P26663; IJXP;
DR InterPro: IPR00345; CytC_hemo_bind.
DR InterPro: IPR01410; DEAD.
DR InterPro: IPR002522; HCV_capsid.
DR InterPro: IPR002521; HCV_core.
DR InterPro: IPR002519; HCV_env.
DR InterPro: IPR002531; HCV_NS1.
DR InterPro: IPR002518; HCV_NS2.
DR InterPro: IPR004109; HCV_NS3.
DR InterPro: IPR006745; HCV_NS4a.
DR InterPro: IPR014490; HCV_NS4b.
DR InterPro: IPR002868; HCV_NS5a.
DR InterPro: IPR002166; HCV_RdRp.
DR InterPro: IPR01650; Helicase_C.
DR InterPro: IPR007095; RNA_pol_PS-ps.
DR InterPro: IPR007094; RNA_pol_PSVit.
DR Pfam: PF01543; HCV_capsid; 1.
DR Pfam: PF01542; HCV_core; 1.
DR Pfam: PF01539; HCV_env; 1.
DR Pfam: PF01560; HCV_NS1; 1.
DR Pfam: PF01538; HCV_NS2; 1.
DR Pfam: PF02907; HCV_NS3; 1.
DR Pfam: PF01006; HCV_NS4a; 1.
DR Pfam: PF01001; HCV_NS4b; 1.
DR Pfam: PF01506; HCV_NS5a; 1.
DR Pfam: PF00271; helicase_C; 1.
DR Pfam: PF00998; Viral_RdRp; 1.

```

```

DR ProDom: PD186062; HCV_NS1; 1.
DR SMART: SMO0487; DEXDC; 1.
DR PROSITE: PS00190; CYTOCHROME_C; 2.
DR PROSITE: PS05057; RDRP_POSITIVE; 1.
DR PROSITE: PS05052; RDRP_VIRAL; 1.
KW ATP-binding; Coat protein; Envelope protein; Glycoprotein; Helicase;
KW Hydrolase; Nonstructural protein; Polyprotein;
KW RNA-directed RNA polymerase; Transferase; Transmembrane.
SQ SEQUENCE 3010 AA; 327177 MW; 47A51DD7678DE62F CR664;

Query Match 95.5%; Score 42; DB 12; Length 3010;
Best Local Similarity 86.9%; Pred. No. 19;
Matches 8; Conservativity 1; Mismatches 6; Indels 0; Gaps 0;

QY 1 LIGCITSL 9
DB 1039 MLCITSL 1047

Search completed: September 29, 2003, 19:08:26
Job time : 32 secs

```